Report

Site Investigation/ Surface Waste Removal

Former Iron Foundry Area South Tacoma Swamp

TIP Management, Inc. Tacoma, Washington

K/J/C 866738 May 1988

USEPA SF 1427584

Kennedy/Jenks/Chilton

Kennedy/Jenks/Chilton

Consulting Engineers

33301 Ninth Avenue South, Suite 100 Federal Way, Washington 98003 206-874-0555 (Seattle) 206-927-8688 (Tacoma)

3 April 1989

Mr. Thomas R. Anderson Tacoma Industrial Properties CS 2259 1123 Port of Tacoma Road Tacoma, WA 98401

Subject: Report on Site Investigation/Surface Waste Removal

Former Iron Foundry Area 5202 S. Proctor Street Tacoma, Washington K/J/C 866738.00

. Dear Mr. Anderson:

Enclosed is the report on our site investigation/surface waste removal at the former iron foundry area, 5202 South Proctor, Tacoma, Washington.

This study has been prepared in accordance with our professional services agreements of 30 September 1986 and 22 October 1987.

Please call us if you have any questions about the information presented in this draft report.

Very truly yours,

KENNEDY/JENKS/CHILTON

Notha Gues

Nathan A. Graves

Manager, Industrial Services

NAG:kae/38

Enclosure

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EXECUTIVE SUMMARY

PROJECT PURPOSE/OBJECTIVES

This report describes environmental studies performed at the Tacoma Industrial Park area located within the boundaries of the South Tacoma Channel Superfund site, Tacoma, Washington. The purpose of this report is to describe field investigations, and evaluate chemical data in the area of a former iron foundry and the surrounding area.

EXTENT OF CHEMICAL COMPOUNDS AT THE SITE

Several documents produced as a result of past site investigations and chemical data generated by Kennedy/Jenks/Chilton were reviewed to assess chemical contamination onsite. Inorganic and organic contaminants were found in site soil, sediment and groundwater at generally low concentrations. Inorganic chemical concentrations were found to be above background soil levels in shallow subsurface soil at one location (lead, copper, cadmium, and zinc) and in four samples of storm drain sediments (lead). It is believed that the inorganics in the soil are limited in extent and relatively immobile, based on nearby soil data and the expected mechanisms that tend to bind metals to soils. A leach test (EP toxicity) was conducted on the sediment sample containing the highest lead concentration and indicated that the lead is expected to be relatively immobile in the sediments.

Organic contaminants detected in site soil and groundwater were generally well below applicable regulatory criteria. The exception was xylenes, which were slightly above WDOE standard cleanup levels at one location near the former solvent shed.

REMOVAL OF WASTE PRODUCTS

Waste materials stored in drums and small containers have been characterized and removed from the site. The wastes were generally composited, based on their characteristics, in order to transport the material in bulk to a permitted disposal facility.

CONCLUSIONS/RECOMMENDATIONS

Contamination at the former iron foundry appears to be limited and is comprised of inorganic compounds in storm drain sediments in one storm drain and in shallow subsurface soil in one location; and organic compounds in shallow soil near the former solvent shed. The removal of the visually stained soils near the solvent shed should be considered, although current risks from this material appear low. WDOE could request the removal of the contaminated soil, based on a recently utilized cleanup level of 200 mg/kg total oil and grease, which the soil is likely to exceed. In addition, the presence of solvents could make removal prudent, due to possible future concerns regarding ingestion or inhalation of the contaminants.

EXECUTIVE SUMMARY, continued

The removal of contaminated storm drain sediments should also be considered, as cleanout of the drains is relatively easy to accomplish. This would eliminate the possibility of contaminated sediments being washed downstream during high rainfall events.

Based on available field data and available data regarding historical site operations, the site appears to have had a minimal impact on overall groundwater quality in the South Tacoma Channel area.

SECTION 1

INTRODUCTION

1.1 PURPOSE

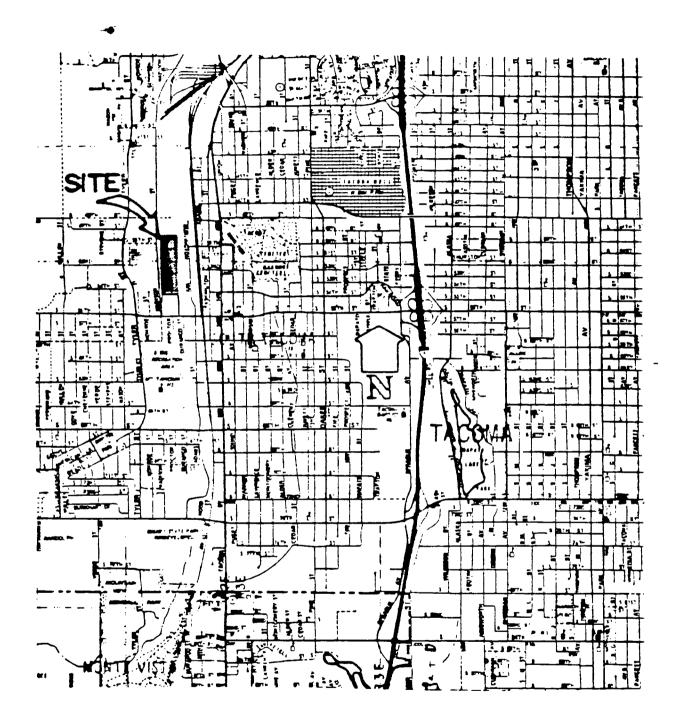
This report represents the culmination of several site assessments by numerous parties related to chemical compounds in soil, sediments and groundwater in the Tacoma Industrial Park area, 5202 S. Proctor Street, Tacoma, Washington. The purpose of this report is to describe field investigations, and evaluate chemical data in the area of a former iron foundry and the surrounding property. An evaluation of chemical concentrations in the vicinity of the adjacent brass foundry was contained in Kennedy/Jenks/Chilton's "Draft Report, Remedial Investigation/Risk Assessment/Feasibility Study, Former Brass Foundry Area", Volumes I and II, and will not be addressed in this report.

Data utilized to form the basis of our site investigation is contained in several documents produced as the result of past site investigations. These documents are listed below:

- o Pierce, D., and Rogers, S., Tacoma Pierce County Health Department, 1982, "South Tacoma Industrial Waste Summary," Tacoma, Washington.
- o Black & Veatch, 1983. Preliminary Site Investigation, South Tacoma Swamp, Tacoma, Washington.
- o Benlab, 1985. "Resource Evaluation of Anderson Enterprises South Tacoma Property."
- o Earth Consultants, 1985. "Report on Soil Sampling and Chemical Testing." Anderson Enterprises Property, Tacoma, WA.
- o Black & Veatch, 1985. Memorandums to EPA/WDOE.
- o Ecology and Environment, 1985. Memorandums and Data Analyses Sheets.

Kennedy/Jenks/Chilton has reviewed these documents and concludes that previous site investigations were conducted in accordance with generally accepted investigation procedures. Based on information contained in these reports, referenced field sampling and analytical techniques should provide adequate data to form the basis for additional investigations. We assume that all data presented in these reports are accurate as stated.

The specific objectives of Kennedy/Jenks/Chilton's study of the iron foundry area were as follows:



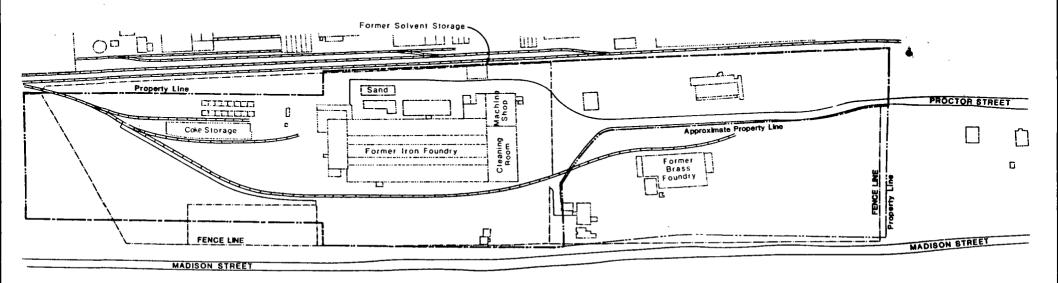
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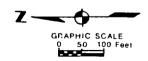
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Tacoma industrial Propertie
Tacoma, Washingto

K/J/C 86673 February 198 Figure 1-





Kennedy/Jenks/Chilton

Tacoma Industrial Properties
Tacoma, Washington
Tacoma Industrial Park Site

K/J/C 866738 February 1988 Figure 1-2

- Assessment of the extent of chemical compounds throughout the area of the former iron foundry based on the review of previous site investigations.
- o Assessment of the extent of chemical compounds in surface and subsurface soils, storm drain sediments and groundwater near the former iron foundry, former solvent storage area and near the reported location of an underground fuel tank.
- o Assessment and evaluation of the environmental risks posed by chemical compounds found onsite.
- o Gross characterization and compatibility testing of materials found in drums/containers onsite in preparation of compositing for disposal offsite at a permitted facility.
- o Detailed analyses of container/drum contents.
- o Resident engineering during the removal of containerized materials for offsite disposal.
- o Development and evaluation of site specific remedial action alternatives, if required, based on the information obtained during the site investigation.

1.2 SITE DESCRIPTION

The former iron foundry plant (subject of this report) is located north of South 56th Street and east of Madison Street, Tacoma, Washington, as shown in Figure 1-1. The plant site is shown in Figure 1-2. The adjacent site, containing the former brass foundry (no longer owned by Tacoma Industrial Properties), is not assessed in this report.

The former iron foundry plant is situated in a localized mixed commercial/industrial setting, and is currently zoned for light industrial (M-2) (Anderson, 1986). The areas north and immediately west of the site are undeveloped, with the areas south and east of the site primarily used for commercial and light industrial activities. The closest area with a large residential population is located approximately one-third of a mile southwest of the site. A few residences are located one-quarter of a mile northeast of the site. The site has been owned by Tacoma Industrial Properties (formerly Anderson Enterprises) since 1974.

The site is located on the floor of an erosional channel that was occupied by a major glacial meltwater stream during the waning stages of the last glaciation of this region, approximately 13,000 years ago. This four-mile long meltwater valley is locally referred to as the "South Tacoma Channel". The channel is incised as much as 150 feet below the rolling upland area that is occupied by residential and commercial districts of Tacoma. The channel

extends north and east of the former iron foundry area to the vicinity of the intersection of South Tacoma Way and Yakima Avenue. South of the study area, the glacial channel widens and opens into a broad glacial outwash plain that is underlain by highly permeable gravel.

The "South Tacoma Swamp" is located immediately north and west of the former iron foundry area. This lowlying portion of the valley floor has been partly filled and drained.

No creeks or streams are present in the vicinity of the property. The headwater area of Flett Creek is presently approximately 6,000 feet south of the site. Flett Creek drains into Chambers Creek and then into Puget Sound.

The Tacoma-Pierce County Health Department has completed a regional geohydrologic study of the Clover/Chambers Creek drainage basin. The results of that study are included in a July 1985 report prepared by Brown and Caldwell entitled, "Clover/Chambers Creek Geohydrologic Study". The Brown and Caldwell report indicates that the site is located in an area where potential contamination of the shallow and deep aquifers is possible. The report also indicates a groundwater flow direction that is generally westward in the vicinity of the site.

Numerous water wells have been drilled in and near the South Tacoma Channel. The well logs on file at the Tacoma-Pierce County Health Department were researched to determine the approximate location and capacity of wells within one mile of the site.

The City of Tacoma owns several municipal wells in a well field located 3,000 to 7,000 feet north-northeast of the site. Some of these wells are capable of yields in excess of 3,000 gpm. This well field does not appear to be downgradient of the site.

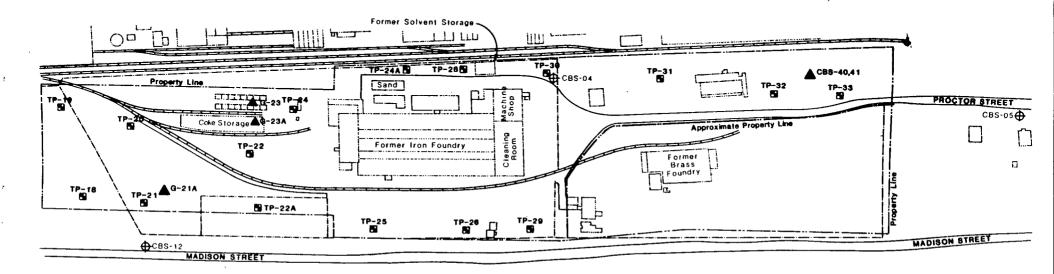
The Town of Fircrest owns three municipal wells located 8,000 to 10,000 feet northwest of the site. Based on information presented in the Brown and Caldwell report, these wells are not located downgradient from the site.

The University Place Water Company owns three public supply wells located approximately 8,000 feet northwest of the site. These wells do not appear to be located downgradient from the site.

Based on available well records, the closest well in the reported downgradient direction is located approximately 2,500 feet west of the site. This private well at 5402 South Mullen Street is 110 feet deep and has a high reported iron content.

1.3 SITE HISTORY

Griffin Wheel, a division of Amsted Industries, bought the former foundry facility and surrounding plant from the American Foundry





LEGEND

- Previously Installed Test Pit
- Approximate Location and Number of EPA Monitoring Well
- A Previous Surface Soil Sample Location

Kennedy/Jenks/Chilton

Tacoma industrial Properties
Tacoma, Washington

Tacoma Industrial Park Site Previous Sampling Locations

> K/J/C 866738 February 1988 Figure 1-3

Company in January, 1897. American Foundry used the plant as a foundry from October, 1890 to 1896.

Griffin Wheel operated two foundries until 1957. The foundry building in the northern portion of Tacoma Industrial Park produced iron wheels until 1957, when the railroads switched to steel wheels. (This building is currently being utilized for other manufacturing activities). The former brass foundry located to the south of the property (not a subject of this report) produced journal bearings until April 1980.

In addition to reviewing past site investigation reports, a review of available aerial photographs of the site was performed. Local aerial photographers were contacted and photographs of the site were obtained. Each photograph was examined for visible indications of contamination (discolored soils) and for indications of activities which may have resulted in site contamination. Dates of the photographs that were reviewed and a discussion of relevant site characteristics and/or activities are provided below.

- o 1946 Iron foundry appears to be operating (wheels are stacked outside of building). Northern Pacific maintenance yards present east of site.
- o 1961, 1969, 1974 No significant change from 1946.
- o 1985 No significant change around iron foundry. Northern Pacific railyards demolished. General Plastics constructed (east of iron foundry).

1.4 HISTORY OF PREVIOUS SITE INVESTIGATIONS

- o October 1982 Black & Veatch performed a preliminary site investigation on the South Tacoma Swamp for the USEPA. The investigation included collection of soil and groundwater samples near the former iron foundry site as shown on Figure 1-3. Applicable sample type and locations relevant to the former brass foundry area include the sampling of groundwater from monitoring wells CBS-04, CBS-05, and CBS-12.
- o November 1982 Black & Veatch resampled the groundwater in wells CBS-04 and CBS-12 as part of the follow-up sampling conducted in the South Tacoma Swamp.
- o February 1985 Benlab and Earth Consultants sampled soil from backhoe pits and assessed the depth and nature of the fill materials in those areas. Figure 1-3 indicates the test pit locations. Benlab analyzed soil samples for total lead and total arsenic.
- o September 1985 Soil samples collected during the February 1985 study were analyzed for EP Toxicity lead and arsenic; and

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additionally for mercury, cadmium, chromium, silver, barium, and selenium on selected samples.

o June 1985 - Black and Veatch and Ecology & Environment resampled the EPA groundwater wells installed in 1982. The sampling effort included wells CBS-04 and CBS-12.

1.5 RESULTS OF PREVIOUS SITE INVESTIGATIONS

Black & Veatch, 1982 and Ecology & Environment, 1985. The results from Black & Veatch's preliminary site investigation (1982) are presented in Tables 1-1 and 1-2. Although methylene chloride, bis(2-ethylhexyl) phthalate and acetone were detected in very small quantities in some samples, they were not shown in the tables. These compounds are common lab contaminants, and may have been detected due to field and laboratory contamination during this investigation. However, quality assurance data is unavailable to confirm this conclusion.

Results of Chemical Concentrations in Groundwater Samples. Groundwater monitoring wells installed by Black & Veatch in 1982 and sampled in 1982 and 1985 include CBS-04, CBS-05, and CBS-12 (Figure 1-3). In relation to the former iron foundry area, CBS-04 and CBS-05 can be considered "upgradient" wells and CBS-12 a "downgradient" well, based on the general northwestern groundwater gradient as reported by Black & Veatch. Priority pollutants detected in upgradient wells CBS-04 and CBS-05 in 1982 include trichloroethylene and zinc. Priority pollutants detected in downgradient wells include pentachlorophenol, 1,1,1-trichlorethane, tetrachlorethylene, zinc, and cyanide. The organic priority pollutants listed above were all detected in trace quantities in the samples, below the quantification limit.

Black and Veatch and Ecology & Environment resampled CBS-04 (upgradient) and CBS-12 (downgradient) in 1985. No organic priority pollutants were detected in either well, including those compounds detected in 1982. In addition, cyanide was not detected in groundwater from well CBS-12. Several priority pollutant metals were detected in groundwater samples from both CBS-04 and CBS-12 (including zinc, as previously detected in 1982); however, most compounds were detected in trace quantities (below quantifiable limit) or concentrations were suspect due to excess holding times prior to sample analyses (Ecology & Environment, Inc., 1985).

Discussion of Chemical Concentrations in Groundwater Samples. The presence of 1,1,1-trichloroethane, tetrachlorethylene and trichloroethylene in 1982 sampling could be a result of documented contamination in the City of Tacoma's South Tacoma well field (Well 12-A) from property in the vicinity of Time Oil Company, northeast of the site, or from documented contamination from the City of Tacoma landfill located west of the site. However, this conclusion cannot be confirmed based on available data. These

compounds were not detected during the 1985 sampling effort at CBS-12 (detection limits in 1985 were also higher). This indicates that groundwater in the vicinity of CBS-12 may not be continuing to be degraded over time by these compounds.

Priority pollutant metals are common elements of the earth's crust and their presence in groundwater is common at low concentrations. The significance of the levels of priority pollutant metals in groundwater was investigated by comparing groundwater concentrations with available drinking water standards as shown on Table 1-2. (Drinking water standards are used because of the use of area groundwater for municipal and private drinking water). For compounds with drinking water standards, all levels detected in groundwater from CBS-04 and CBS-12 are below drinking water standards.

The source of cyanide, which was detected at CBS-12 during one sampling event in 1982, is unknown. However, cyanide was not detected in CBS-12 during a second sampling event in 1982. The levels of cyanide detected in 1982 (17 ppb) is below the level of public health significance (200 ppb) as reported by the U.S. EPA (EPA, 1980). In addition, cyanide was not detected in CBS-12 in 1985.

Chemical Concentrations in Soil Samples. Chemical compounds detected in soil samples from the installation of Black & Veatch's monitoring wells and the collection of surface soil samples in 1982 are summarized in Table 1-3. The inorganic priority pollutants detected in soils from the monitor wells and the one surface soil sample (CBS-40) were consistently low in concentration and well within typical ranges of these metals in soils. However, some metals in soil just below the surface at one location (CBS-41) were high, namely cadmium (1.7 ppm), lead (4300 ppm), copper (2800 ppm) and zinc (1800 ppm). Inspection of the logs for test pits installed by Earth Consultants, 1985, in the area of the elevated metals, showed a material described as "dark sand with metallic luster" at depths of 1-1/2 feet and 8-9 feet in two test pits (TP-32 and TP-33). The high metals concentrations could be associated with this material. EP toxicity analyses both above and below the metallic sand resulted in non-detectable quantities of leachable lead and arsenic, indicating the reduced mobility for these compounds. In addition, soil at only one location exhibited these high levels of metals, indicating limited contamination.

Organic priority pollutants were not detected in soil from monitor well CBS-04. Trichlorofluoromethane was only detected in soil from CBS-12 at 13.5 feet at trace concentrations (6.0 ppb). Trichloroethylene was only detected at trace concentrations (32 ppb) in the surface soil sample (CBS-40). Organic compounds which were also detected at trace concentrations but which were below the specified detection limit were fluoranthene, chrysene, phenanthrene (CBS-41) and pyrene (CBS-40, CBS-41).

TABLE 1-1
SUMMARY OF PRIORITY POLLUTANT ANALYSIS OF GROUNDWATER SAMPLES (a)
TIP MANAGEMENT - IRON FOUNDRY SITE

Sample Location Well Number	Upgradient CBS-04			nt-Offsite S-05		radient S-12	Drinking Water Standards	
Date	10/20/82	11/11/82	10/19/82	11/04/82	10/21/82	11/10/82	(b)	
Pentachlorophenol	ND	ND	ND	ND	3.0 (c)	ND		
1,1,1-Trichloroethane	ND	ND	ND	ND	6.7 (c)	9.2 (c)	200	
Tetrachloroethylene	ND	ND	ND	ND	3.5 (c)	3.0 (c)	0	
Trichloroethylene	2.1 (c)	ND	ND	ND	ND	ND	5.0	
Zinc	ND	160	ND	650	32	100	5000	
Cyanide	ND	ND	ND	ND	17	ND	200 (d)	

Notes:

All units are ppb.

a - Black & Veatch, June, 1983.

b - National Primary & Secondary Drinking Water Regulations, 52 FR 41546 and 51 FR 11412.

c - Below quantitation limit.

d - U.S. Public Health Service, 1968.

ND - Not detected.

All other priority pollutants were not detected, except for methylene chloride,

bis (2-ethylhexyl) phthalate, and acetone, which are suspected lab contaminants.

TABLE 1-2 SUMMARY OF INORGANIC PRIORITY POLLUTANT ANALYSES OF GROUNDWATER SAMPLES (a) TIP MANAGEMENT - IRON FOUNDRY SITE

Sample Location Well Number Date	Upgradient CBS-04 06/20/85	Downgradient CBS-12 06/20/85	Drinking Water Standards (b)
Antimony	52 (c)	70	NA NA
Barium (d)	26	11 (c)	1000
Cadmium	4	3.6 (c)	10
Copper	15 (c)	20	1000
Lead	2.4 (c,e)	2.4 (c,e)	50
Mercury	0.20 (e)	0.20 (e)	2
Silver	9.6 (c,e)	9.6 (c)	50
Thallium	2.6 (c,e)	2.6 (c,e)	NA
Zinc	128 (e)	37 (e)	5000

Notes:

All data reported in ppb.

All other inorganic priority pollutants were not detected. NA - None Available

a - Ecology and Environment, Inc., 1985.

- b National Primary & Secondary Drinking Water Regulations, 52 FR 41546 and 51 FR 11412.
- c Material not detected number is estimated sample quantitation limit.
- d Not a priority pollutant, however barium is a regulated drinking water contaminant.
- e Value is an estimated quantity because quality control criteria were not met.

TABLE 1-3

SUMMARY OF PRIORITY POLLUTANT ANALYSES OF SOIL SAMPLES (a)
TIP MANAGEMENT - IRON FOUNDRY SITE

Sample Location	СВ	S-04	СВ	S-12	CBS-40	CBS-41	Typical Soil	EPC's
Date Depth (ft)	10/11/82 17.8	10/11/82 37.8	10/13/82 13.5	10/13/82 28.5	10/18/82 Surface	10/18/82	Range (b)	(c)
Organics								
Fluoranthene	ND	ND	ND	ND	ND	0.4 (d)		
Chrysene	ND	ND	ND	ND	ND	0.4 (d)		
Phenanthrene	ND	ND	ND	ND	ND	0.4 (d)		
Pyrene	ND	ND	l ND	l ND	0.4 (d)	0.4 (d)		
Trichlorofluoromethane	ND	ND ND	0.006	ND	ND`	ND`		15.500
Trichloroethylene	ND	ND	ND	ND	0.032	ND		1.477
Inorganics	; i							
Arsenic	ND ND	ND	1.3	1.1	2.3	6.7	1-50	
Cadmium	0.2	0.2	ND	ND '	0.2	1.7	0.01-0.7	
Chromium	1.3	2.2	l ND	ND	ND	ND	1-1000	
Copper	ND ND	ND	ND	ND	20	2800	2-100	
Lead	1.0	1.3	3.9	1.3	54	4300	2-200	
Nickel	ND	4.6	ND	ND	ND	ND	5-500	
Zinc	1.9	5.0	6.2	2.9	11	1800	10-300	
Zinc	1.9	5.0	6.2	2.9	11	1800	10-300	

Notes:

- All units are ppm. ND Not Detected.
- All other priority pollutants were not detected, except for methylene chloride and acetone, which are suspected lab contaminants.
- a Black & Veatch, June 1983.
- b Lindsey, W.L., Chemical Equilibria in Soil, Wiley & Sons, New York, 1979.
- c Estimated Permissible Concentrations, US EPA, "Multimedia Environmental Goals for Environmental Assessment", Vols. I, II, EPA 600/7-80-041.
- d Detected below, but greater than one half the specified detection limit.

Benlab, 1985

Benlah only analyzed one subsurface and one surface sample for total lead and arsenic near the iron foundry. Soil from a depth of 4.5-6 feet in TP-25 contained lead at 477 ppm and arsenic at 0.04 ppm. The arsenic level is below its typical soil range, while the lead exceeds its typical range in soil. EP toxicity analyses of the same sample (Table 1-5) resulted in non-detectable lead, indicating its immobility in the soil system.

Earth Consultants, 1985

EP toxicity analyses of soil samples were performed to assess chemical concentrations at the site compared with one regulatory criteria. The EP toxicity analysis is a test that indicates the relative "leachability" of a compound under acidic conditions (simulating sanitary landfill leachate). This test does not generally simulate leachability in the normal environment, but the test is used to assess proper disposal practices for wastes.

The results of EP toxicity analyses, on soil, performed by Earth Consultants (1985) are shown in Table 1-4 (lead and arsenic), and in Table 1-5 (lead, arsenic, mercury, cadmium, chromium, silver, barium and selenium). Arsenic, mercury, cadmium, chromium, silver, barium and selenium were not detected. Lead was detected in four samples at trace concentrations. None of the samples had concentrations exceeding the State of Washington's dangerous waste criteria for lead.

SUMMARY

Trace levels of organic priority pollutants were detected in site soil and groundwater in two prior studies at the iron foundry plant site. Analyses for metals in site groundwater detected zinc in both sampling rounds, with additional metals detected in 1985 sampling event. All of these metals concentrations were well below drinking water criteria. Metals were also detected in site soil, with concentrations within typical soil levels, except for one location. One shallow soil sample contained elevated levels of cadmium, copper, lead and zinc. However, these levels appear to be limited in extent and exhibit limited mobility. EP toxicity analyses of site soil (lead and arsenic) for comparison to Dangerous Waste criteria showed the soil is not a Dangerous Waste. Arsenic was not detected, using EP toxicity test, while lead was detected in trace concentrations at four locations.

It appears that inorganic site contamination is confined to one localized area around CBS-41, while organic contamination appears to be at trace levels in both soil and groundwater at CBS-12, in soil at CBS-40, and groundwater at CBS-04. However, an onsite source of groundwater contamination cannot be confirmed, and contaminants may be present as a result of several documented sources nearby the site.

TABLE 1-4

SUMMARY OF EP TOXICITY LEAD AND ARSENIC IN SOIL SAMPLES (a)

TIP MANAGEMENT - IRON FOUNDRY SITE

Test Pit Depth (ft)	TP-18 0-2	 TP-18 1-1.5 	 TP-18 8-10 	TP-19 0-1.5	TP-19 8-9	 TP-21 * 0-1 	TP-21 7-9	TP-22 10-12	 TP-22A 2-4	 TP-24 0-1	TP-24 7-8	 TP-24A 0-1 	 Dangerous Waste Criteria (b)
Lead Arsenic	<0.1 <0.1	 <0.1 <0.1	 <0.1 <0.1	0.2 <0.1	<0.1 <0.1	 <0.1 <0.1	<0.1 <0.1	<0.1 <0.1	 <0.1 <0.1	 <0.1 <0.1	<0.1 <0.1	 <0.1 <0.1	 5-500 5-500
Test Pit						1	TP-28	TP-28	:	 TP-29	TP-29		Dangerous Waste
Depth (ft) Lead Arsenic	<0.1 <0.1	1-3 <0.1 <0.1	4.5-6 <0.1 <0.1	13-14 	3-5 0.3 <0.1	16-17 <0.1 <0.1	0-0.5	2-3 <0.1 <0.1	6-7 	2-3 	2-4 <0.1 <0.1	5-7 <0.1 <0.1	Criteria (b)
· 		i 	i	i	i	l		l	 	!		<u></u>	1
Test Pit Depth (ft) 		TP-30 0-1	TP-30 7-8 	TP-31 0-1 	TP-31 6-7	TP-32 1-1.5 	TP-32 2-3	TP-33 2-4 	TP-33 9-11 	G-21A Surface 		•	Dangerous Waste Criteria (b)
Lead Arsenic	<0.1 <0.1	0.4	<0.1 <0.1	 <0.1 <0.1	<0.1 <0.1	<0.1 <0.1 <0.1	<0.1 <0.1	<0.1 <0.1	<0.1 <0.1	<0.1 <0.1 	<0.1 <0.1	<0.1 <0.1 <0.1	5-500 5-500

Notes:

- a Earth Consultants, 1985.
- b Dangerous Waste Criteria, Chapter 173-303 WAC, amended June 1987.
- * Selected for additional EP Toxicity metals analyses.

TABLE 1-5 SUMMARY OF EP TOXICITY ANALYSES OF SOIL SAMPLES (a) TIP MANAGEMENT - IRON FOUNDRY SITE

Test Pit Number Depth (ft)	TP-21 0-1	TP-25 1-3	TP-25 4.5-6	TP-25 13-14	Dangerous Waste Criteria (b)
Lead	<0.1	<0.1	<0.1	<0.1	5-500
Arsenic	<0.1	<0.1	<0.1	<0.1	5-500
Mercury	<0.01	<0.01	<0.01	<0.01	0.2-20
Cadmium	<0.01	<0.01	<0.01	<0.01	1-100
Chromium	<0.01	<0.01	<0.01	<0.01	5-500
Silver	<0.01	<0.01	<0.01	<0.01	5-500
Barium	<1	<1	<1	<1	100-10,000
Selenium	<0.1	<0.1	<0.1	<0.1	1-100

Notes:

- All data reported in ppm. a Earth Consultants, 1985. b Dangerous Waste Criteria, Chapter 173-303 WAC, amended June 1987.

SECTION 2

SITE INVESTIGATION

2.1 PROJECT OBJECTIVES

The purpose of Kennedy/Jenks/Chilton's additional site investigation of the former iron foundry and the surrounding plant was to gather the additional data necessary to assess the risks posed by chemical compounds on the site, as well as to form the basis for evaluating remedial action alternatives, if required. To gather this data, several evaluations or investigations were conducted as listed below:

- Review of past site investigations.
- Development and performance of soil, sediment, and groundwater investigations to fill gaps in the database.
- Performance of containerized material investigations to establish material characteristics for offsite disposal.

The results of the investigations described above were then used to fulfill the overall objectives of the site investigation, which include:

- Characterize the areal extent of chemical compounds in various site media.
- Identify possible public health and/or environmental exposures posed by chemical compounds at the site,
- Removal of drummed materials for offsite disposal, and
- Identify remedial action needs.

2.2 SITE INSPECTION

A detailed walkthrough of the former iron foundry plant was performed to observe possible areas of visible surface soil contamination and to identify container storage areas. The results of our inspection and review of available site data were used to select appropriate sampling locations as needed to further identify the presence and extent of chemical contamination at the site.

Stained surface soils were present near a former solvent shed east of the foundry, and near a reported transformer oil storage area north of the foundry. Drums were sighted in several locations throughout the site. Numerous small containers were present within the former brass foundry building southwest of the iron foundry.

parper PCBs?

2.3 FIELD INVESTIGATION PROCEDURES

2.3.1 Surface Soil Sampling

On 2 October 1986, samples of surface soils were collected in two locations near the former iron foundry: one near a reported former transformer oil storage tank and one by the former solvent storage area. Samples were collected using clean plastic scoops and were immediately transferred into clean glass jars and cooled, prior to shipment to Kennedy/Jenks/Chilton Laboratory Division under chain-of-custody procedures.

2.3.2 Storm Drain Sampling

On 9 and 10 October 1986, samples were collected from seven storm drains around the former iron foundry. Two of the drains (SD-2 and SD-5) did not contain any sediments, so grab water samples were collected. Sediments were collected from the other five drains. The sediment samples were collected using clean plastic scoops and immediately transferred into clean glass jars. The water was collected directly into clean glass jars. All storm drain samples were stored on ice and shipped to our Laboratory Division under chain-of-custory procedures.

2.3.3 Boring/Monitor Well Installations

On 11 November 1987, four soil borings and one monitoring well were completed at locations near the former solvent shed and the reported location of an underground fuel storage tank as shown on Figure 2-1. The drilling work was contracted to F.L.D. Industries located in Everett, Washington. A Kennedy/Jenks/Chilton engineer logged the boring operation and installation of the monitoring well. Well logs are presented in Appendix A. A truck-mounted drilling rig was used. Before work at the site began and between each boring, the drilling equipment was steam cleaned to reduce the likelihood of cross-contaminating the borings or samples. Soil samples were taken with a standard split-barrel sampler.

Borings B-1, B-2, and B-3 were drilled in areas of stained soil in the solvent storage area. The borings were terminated below the visually contaminated soil. Based on visual observations during these borings, an estimate of the volume of stained soil in this area was calculated to be approximately 2,000 cubic feet. Boring B-4 was positioned near the reported location of an underground gasoline tank. The tank bottom was measured to be 8 feet below the ground surface by using a weighted tape measure. The tank contained approximately 3-3/4 inches of liquid which had an odor of gasoline. The boring placed adjacent to the tank (B-4) was terminated 2-1/2 feet below the bottom of the tank. Gasoline odors were not detected in any of the soil samples or drilling cuttings. Monitoring well MW-5 was placed at a point

expected to be downgradient from the solvent storage area. MW-5 was constructed with 2-inch diameter PVC screen and casing. The screen was machine slotted and 20 feet long. A filter pack was placed surrounding the screen, and a bentonite seal was placed above the filter pack. The top of the well was sealed with a water-tight locking cap and covered with a traffic-rated monument cover. Details of the construction of this monitoring well are shown on the well log in Appendix A.

2.4 HYDROGEOLOGIC CONDITIONS

Subsurface soil conditions beneath the site were explored by drilling three shallow borings and one deeper boring. The shallow borings encountered sandy gravel fill extending to the base of the borings at approximately five feet. The deeper boring encountered fine sand with less than five percent gravel extending to the base of the boring at approximately eleven feet. In addition, a monitoring well was installed in a fifth boring. This boring encountered gravelly sand fill mixed with debris overlying gravelly sand which extended to approximately 32 feet. The gravelly sand unit was deposited by glacial meltwater during formation of the South Tacoma Channel. The base of the gravelly sand unit was not reached.

Groundwater conditions at the site were explored by installing one monitor well and utilizing monitor wells installed near the site by EPA in 1983. The water table elevations in the three EPA wells were measured on September 26, 1986. Groundwater occurs in an unconfined condition within the gravelly sand unit encountered in the wells. Based on water table elevation measurements, the groundwater flow direction beneath the site is generally north-westerly.

2.5 CHEMICAL CONCENTRATIONS/DATA EVALUATION

2.5.1 Surface and Subsurface Soils

Table 2-1 presents a summary of the chemical analyses results of the five surface and subsurface soil samples. Based on results of previous analyses and past use of the area, the only chemicals analyzed in these samples were organic priority pollutant compounds. Results of analyses showed only detectable levels of volatile organic compounds. The shallow soil sample (12-18 inches) from the area of the former solvent shed (GS-2) contained trace concentrations of 1,1,1-trichloroethane (0.55 ppm), tetrachloroethylene (0.3 ppm) and xylenes (5 ppm). To assess the extent of these compounds in subsurface soil near this location, soil from boring B-2 (depth of 5-6.5 feet) was analyzed and found to contain only 1,1,2-trichloro-1,2,2-trifluoroethane at 0.007 ppm. Comparison of site concentrations of these compounds with estimated permissible concentrations (EPCs), calculated by the EPA based on health effects in soil, showed site concentrations

TABLE 2-1

SUMMARY OF CHEMICAL ANALYSES OF SOIL SAMPLES (a)
TIP MANAGEMENT - IRON FOUNDRY SITE

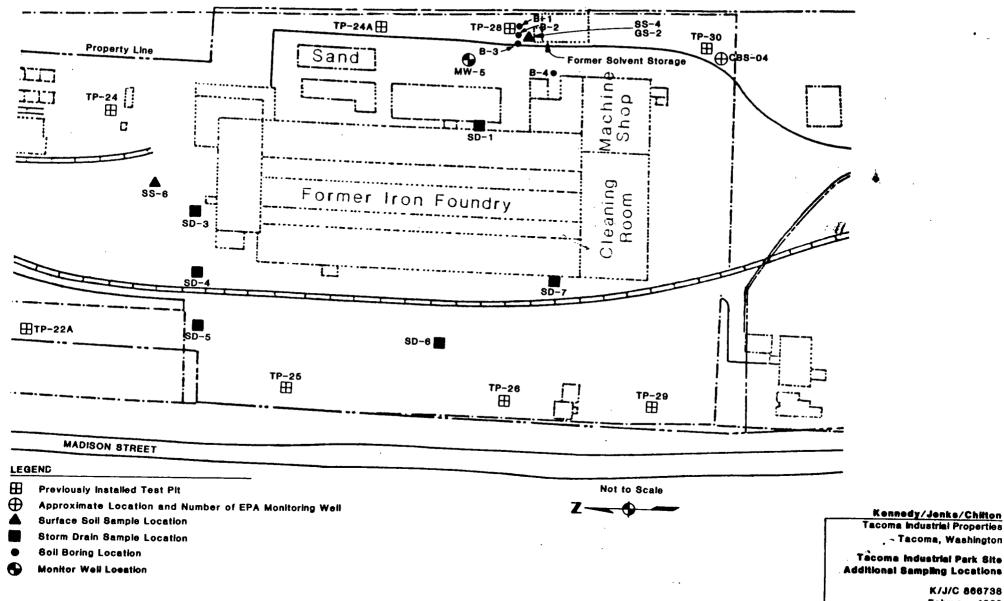
Sample Location Depth (ft) Date	GS-2 1-1.5 10/2/86		SS-6 Surface 10/17/86	B-2 5-6.5 11/12/87	B-4 9-10.5 11/12/87	EPC's (b)		OE anup Policy Protection
Volatile Organics 1,1,1-Trichloroethane Tetrachloroethylene 1,1,2-Trichloro-1,2,2- trifluoroethane Xylenes	(c) 0.55 0.3 <0.1	NA	NA	(d) <0.005 <0.005 0.007 <0.005	NA	1.477 1.849 20.976	2.0 0.7 N/A 4.4	20.0 7.0 N/A 44.0
EPA Method 8270	ND (e)	ND (f)	NA	NA	NA			
Purgeable Aromatics (g)	NA	NA	NA	NA	<0.001			
PCBs	NA	NA	<0.001	NA	NA			

Notes: ND - Not Detected. NA - Not Analyzed For. N/A - None Available.

All units are ppm.

a - Kennedy/Jenks/Chilton, 1986-87.

- b Estimated Permissible Concentrations, US EPA, "Multimedia Environmental Goals for Environmental Assessment," Vols. I, II.
- c No other volatile organics were detected. Detection limits ranged from 0.1 to 2.0 mg/Kg
- d No other volatile organics were detected. Detection limits ranged from 0.005 to 0.03 mg/Kg.
- e Compound detection limits ranged from 0.2 to 1.0 mg/Kg.
- f Compound detection limits ranged from 2.0 to 10.0 mg/Kg.
 g Purgeable aromatic compounds were not detected. All of these compounds had a detection limit of 0.001 mg/Kg.



Kennedy/Jenke/Chitton Tacoma Industrial Properties

Tacoma Industrial Park Site Additional Sampling Locations

> K/J/C 866738 February 1988 Figure 2-1

were many orders of magnitude less than the EPCs. EPCs were used due to the lack of available data regarding acceptable levels of organic compounds in soils. Calculations utilized to develop EPCs are shown in Appendix B.

The Washington Department of Ecology (Ecology) has presented a Final Cleanup Policy, which pertains to cleanup standards for chemical coumpounds in soil, groundwater, surface water, and air. The cleanup levels for soil are primarily based on multiplicative factors of water quality criteria or drinking water standards. The standard/background cleanup levels are ten times the water standard, while the protection cleanup level is 100 times the standard. The standard cleanup level essentially specifies the level at which a site is considered "clean." Protection cleanup levels are applicable only with long-term monitoring to assess that drinking water is not being degraded. Drinking water standards are available for 1,1,1-trichloroethane and xylenes. Since a drinking water standard is not available for tetrachloroethylene, an acceptable water concentration was calculated for tetrachloroethylene, (see Appendix B) using an acceptable increased cancer risk level of 10-5 and its carcinogenic potency slope. Comparison with site soil concentrations show that xylenes are slightly above the standard level, and none of the compounds are above the protection level. However, due to the limited area of contamination, we suspect that chemicals in soil will provide a minimal risk to groundwater, due to the small area of contamination and the high productivity of the underlying aquifer (thus diluting the contaminants). Ecology has also recently used a level of 200 mg/kg total oil and grease as a cleanup level on contaminated sites. It is likely that all the visually contaminated soil in the solvent shed area contains in excess of this level; therefore, Ecology could request that the material be removed for offsite disposal. This action may be prudent, considering the presence of solvents in the soil and possible future concerns regarding ingestion or inhalation of these compounds. However, current risks posed by these materials appear low.

Purgeable aromatic hydrocarbons (benzene compounds, toluene, xylenes) were not detected in soil from boring B-4, adjacent to an underground fuel storage tank. PCBs (polychlorinated biphenyls) were not detected in the surface soil sample (SS-6) taken near a reported transformer storage area. Analysis by EPA Method 8270 did not detect acid/base/neutral extractable organics in soil from GS-2 or SS-4 (from solvent storage area).

2.5.2 Storm Drain Sediments

Seven storm drain locations were sampled for lead and total petroleum hydrocarbon (TPH) analyses. The results of these analyses are presented in Table 2-2. Total lead concentrations in the sediments of four of the storm drains exceeded typical soil

TABLE 2-2

SUMMARY OF CHEMICAL ANALYSES OF STORM DRAIN SAMPLES (a)
TIP MANAGEMENT - IRON FOUNDRY SITE

Sample Location Type Date	SD-1 Sed. 10/9/86	SD-2 Water 10/9/86	SD-3 Sed. 10/10/86	SD-4 Sed. 10/9/86	SD-5 Water 10/9/86	SD-6 Sed. 10/9/86	SD-7 Sed. 10/9/86	Typical Soil Levels (b)
Total Lead (mg/Kg)	320	NA	530	96	NA	1200	310	2-200
Total Lead (mg/L)	NA	9.0	NA	NA	1.6	NA	NA	
EP Toxicity (mg/L) Lead	NA	NA	NA .	NA	NA	<0.5	NA	
TPH (mg/Kg) (c)	NA	NA	0 5	NA	NA	NA	i	
as Diesel Fuel as Kerosene			8.5		,		61	

Notes:

a - Kennedy/Jenks/Chilton, 1986.

b - Lindsey, W.L., Chemical Equilibria in Soil, Wiley & Sons, New York, 1979.

c - Total Petroleum Hydrocarbons.

NA - Not analyzed for.

levels for lead. One sediment sample was within the typical range. Lead concentrations varied from 96 ppm to 1200 ppm in sediments. The location with the highest lead concentration (SD-6) was further analyzed for EP Toxicity lead, to assess if the lead would leach from the sediments. The results indicate that the lead should be relatively immobile in the sediments. Two of the storm drain locations (SD-2, SD-5) did not contain sediments, therefore, water samples collected from these locations were analyzed for total lead, with results of 9.0 ppm and 1.6 ppm, respectively. These results probably do not indicate high levels of lead in the water itself, since samples contained high levels of solids. For comparison purposes, detected levels are better compared with typical soil levels in the environment.

Sediments at two locations were analyzed for TPH, SD-3 and SD-7. Diesel fuel was detected at SD-3, while kerosene was detected at SD-7, both at low concentrations.

2.5.3 Groundwater

In addition to re-sampling the monitor wells installed by Black & Veatch (CBS-04,-05,-12) and analyzing for copper and lead, one new well (MW-5) was installed and sampled solely for volatile organic priority pollutants in the downgradient direction from the former solvent shed. The results from these analyses are presented in Table 2-3.

Copper was not detected in either the dissolved or particulate state in any of the Black & Veatch wells. Total lead was detected in all three wells, with dissolved lead only present in CBS-04. All of the lead concentrations detected are below the drinking water criteria of 0.05 mg/l; however, the total lead concentration at CBS-12 is just above the proposed recommended maximum contaminant level (RMCL) of 0.02 mg/l.

Water from MW-5 was analyzed for volatile organic compounds to assess if contaminants from the former solvent shed had entered the groundwater. The only volatiles detected were trichlorofluoromethane (0.2 ppm) and 1,1,1-trichloroethane (0.034 ppm). Drinking water criteria for 1,1,1-trichloroethane (1,1,1-TCA) was not exceeded. Trichlorofluoromethane was not detected in soil samples near this location and no regulatory criteria is available for this compound, so site concentrations were compared with the EPA's estimated permissible concentrations (EPCs). Site concentrations for both compounds were several orders of magnitude less than their respective EPCs.

Conductivity, temperature, pH, and oxygen-reduction potential (ORP) were measured in CBS-04,-05,-12 to help characterize the groundwater. The pH and temperature were in the normal ranges. In addition, conductivity measurements were within the typical range of 50-1500 umhos/cm for potable water.

TABLE 2-3 COMPARISON OF GROUNDWATER CONTAMINANTS WITH REGULATORY CRITERIA (a) TIP MANAGEMENT - IRON FOUNDRY SITE

Sample Location Date	MW-5 11/12/87	CBS-04 9/26/86	CBS-05 9/26/86	CBS-12 9/26/86	Drinking Water Standards (b)	Estimated Permissible Concentrations (c)
pH	NA	7.39	7.46	7.17	6.5-8.5	
ORP (mv) (d)	NA	443	273	460		
Conductivity (umhos)	NA	189	115	185		
Temperature (cel)	NA	10.3	11.5	12.0		
Inorganics	<u> </u>					
Total Copper	l na	<0.01	<0.01	<0.01	1.0	
Dissolved Copper	l NA	<0.01	<0.01	<0.01		
Total Lead	NA NA	0.02	0.02	0.03	0.05 (0.02 e)	
Dissolved Lead	NA	0.01	<0.01	<0.01	•	
Volatile Organics (f)						
Trichlorofluoromethane	0.2	ND	ND	ND	·	77.300
1,1,1-Trichloroethane	0.034	l nd	l ND	ND	0.2	7.383

Notes:

- All units are ppm unless otherwise indicated.
- a Kennedy/Jenks/Chilton, 1986-87.
- b National Primary & Secondary Drinking Water Regulations, 52 FR 41546 and 51 FR 11412. c US EPA, "Multimedia Environmental Goals for Environmental Assessment," Vols. I & II, EPA 600/7-80-041.
- d Reported relative to standard hydrogen electrode.
- e Proposed RMCL, 50 FR 46936, November 13, 1985.
- f No other volatiles detected. Detection limits ranged from 0.01 to 0.06 mg/L.
- NA Not Analyzed

2.5.4 Public Health and Environmental Concerns

Data generated by Kennedy/Jenks/Chilton and others during numerous investigations of the former iron foundry plant site have shown that although some contaminants are present in site soil and groundwater, concentrations are generally below applicable criteria and/or within typical background levels. The exception to this is the soil around CBS-41, which contained elevated heavy metal contamination, and the storm drain samples, which contained elevated lead concentrations.

However, the elevated metals at CBS-41 are not at the surface, reducing the possibility of impact to nearby human and terrestrial life. The storm drain sediments appear limited in extent and should not pose a significant risk of direct contact to biota. High rainfall events could promote washing of the sediments into the Tacoma storm drain system; however, impacts due to the detected levels of lead should be minimal when washed downstream with large volumes of water.

Although existing data does not indicate that the soil/sediments must be removed, cleanout of storm drain sediments is relatively easy to accomplish.

The solvent concentrations present in soil and groundwater near the former solvent shed appear to pose a minimal risk to human health and the environment. The area of contamination is small and appears to be primarily below the ground surface. This reduces the risk to both human and terrestrial life of exposure to organic contaminants at this location.

SECTION 3

CHARACTERIZATION AND DISPOSAL OF WASTE PRODUCTS

3.1 DRUM/CONTAINER SAMPLING/CHARACTERIZATION

In 1987, Kennedy/Jenks/Chilton and Crowley Environmental personnel performed a site reconnaissance of the site for drummed materials. The plant was divided into seven sections to organize the search. Each section was cleared of brush so that drums would not remain hidden. As drums were located, they were transported to the staging area. The staging area was located on a concrete pad near the former iron foundry building. Prior to placing drums in this area, the pad was covered by visqueen and a berm built up around the edges so that any spillage would be contained. Drums in poor condition were placed in overpack drums to reduce the chance of accidental spillage. A total of twenty-four drums were located.

After the drums were staged, Crowley personnel donned personal protective equipment for opening and sampling the drums. Liquids were sampled using dedicated glass rods and solids were sampled using dedicated plastic scoops/spoons. All samples were collected directly into clean glass jars. Observations regarding the visual characteristics of the material, quantity in the drum, labeling on the drum, etc., were noted at ths time.

The individual samples were then characterized using pH, OVA, and the presence of water and chlorinated hydrocarbons. Based on these results, it was determined that all of the liquids should be compatible, except for one acid (pH of less than 1). A composite sample of the liquids was collected by removing approximately the same amount from each individual sample and collecting it in a jar while monitoring the temperature. After compositing the liquids, a small amount of the acid was added to a small amount of the composite in a separate jar to see if there would be a reaction. A large temperature increase showed that the acid should not be combined with the composite for disposal.

A composite of the solid materials was not collected, as it was necessary to dispose of the solids within industrial drums. The composite liquid sample, the acid, and the solids samples were then taken to a local laboratory under chain-of-custody procedures for the analyses required for acceptance by the disposal facility.

3.2 DISPOSAL OF DRUMMED/CONTAINERIZED MATERIAL

866738.00

After receiving the results of the chemical analyses, the containerized wastes were scheduled for removal to an appropriate disposal facility.

Kennedy/Jenks/Chilton

In March, 1988, the removal of the drummed materials in the area of the former iron foundry began. Three drums of solidified black tar and one drum of acid were placed in overpacks by Crowley Environmental and transported to Chempro's Lucile Street facility under manifest number 07151 (copies of all manifests are included in Appendix D). The drums of liquid were opened and vacuumed into a small tank truck. The liquid drums were then triple-rinsed, with the rinsewaters also going into the tank truck. The liquids were taken to Chempro's Lucile Street facility under manifest number 07105.

Other drums of non-dangerous waste were removed to Chempro's Lucile Street facility under manifest number 07152.

SECTION 4

CONCLUSIONS AND RECOMMENDATIONS

CONCLUSIONS

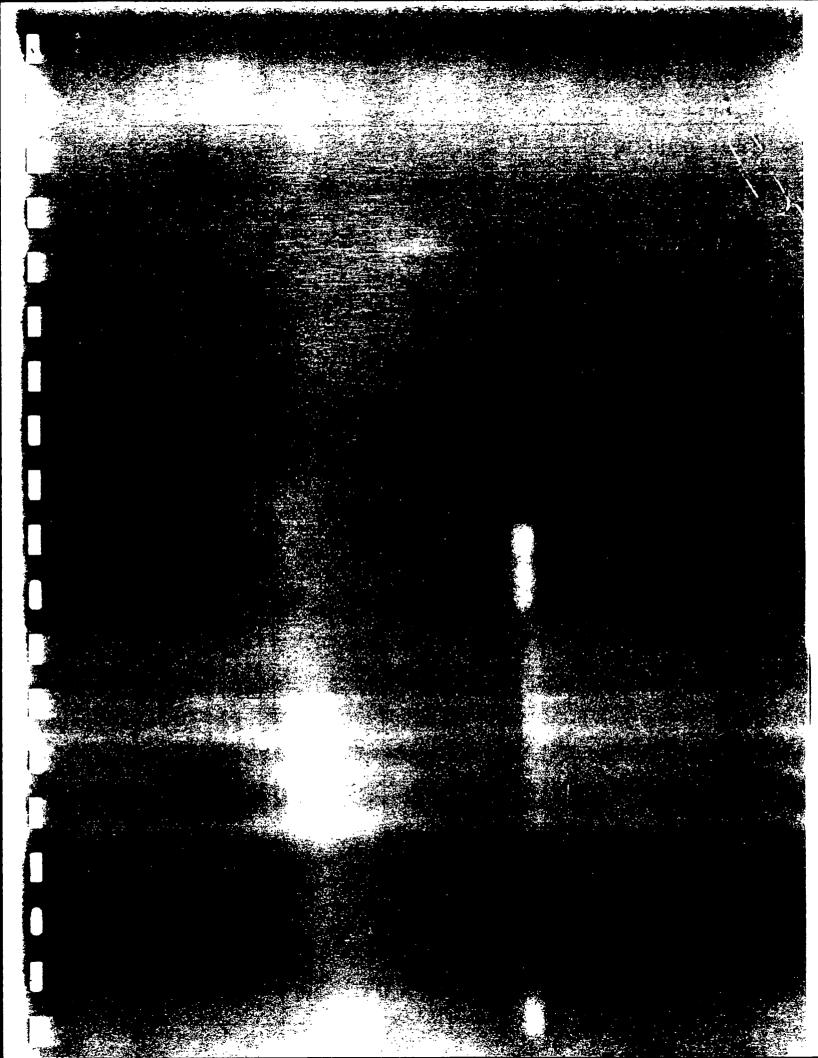
- 1. Based on available information, elevated metals concentrations are present at the former iron foundry plant site in two areas: shallow subsurface soil at CBS-41, and storm drain sediment. However, due to the suspected limited volume of soil with elevated levels of metals and since potential exposure to these metals appears limited, this soil appears to pose a minimal risk to the environment.
- 2. Based on available field data and available information regarding historical operations conducted onsite, the site appears to have had a minimal impact on overall groundwater quality in the South Tacoma Channel area.
- 3. Trace concentrations of four volatile organics (1,1,1-trichloroethane, tetrachloroethylene, xylenes, 1,1,2-trichloro-1,2,2-trifluoroethane) were found in shallow soil near the former solvent shed. All levels were below their EPCs, while levels of xylenes were above WDOE Final Cleanup Policy Standard/Background Cleanup levels but below the Protection Cleanup level.
- 4. Purgeable aromatic hydrocarbons were not detected in soil adjacent to an underground fuel tank.
- 5. PCBs were not detected in soil in the vicinity of a reported former transformer oil storage tank.
- 6. Trace concentrations of lead, below drinking water criteria, were detected in groundwater from the EPA wells. Trace concentrations of two volatile organics (trichlorofluoromethane, 1,1,1-trichloroethane), below applicable criteria, were detected in MW-5, near the former solvent shed.
- 7. Waste materials contained in drums and small containers, as well as building debris, have been removed from the site. A few additional drums of non-dangerous waste remain onsite.

RECOMMENDATIONS

1. Consider removal of visually stained soil from the area of the former solvent shed. Although concentrations of priority pollutants appear to pose a minimal risk to the environment, these soils appear to exceed background cleanup levels and oily soil cleanup quidelines of the Washington Department of Ecology. A

preliminary cost estimate range for removal, transportation, and disposal is \$5,000 to \$10,000, based on 75 yd 3 of material and assuming the material can be disposed of in a construction debris landfill.

2. Evaluate the cost/benefit of removing the storm drain sediments versus leaving them onsite.



Figure

	igure						retinedy denies of inter
PROJ	0.0		a Industrial Proper	ties			BORING B-1
JOB	Details of Boring and Well Construction						
SAMPLE NUMBER	3 Well Construction		DEPTH IN FEET	nscs	SYMBOLS	SURFACE ELEVATION: DRILLING METHODS: P-50 truck-mounted, hollow stem auger. SAMPLING METHODS: Split barrel sampler. DRILLER: Rick Seiler (F.L.D.)	
				-1 -2 -3 -4 -5 -6			0 0' to 6½' Sandy gravel fill, stained dark 0' to approx. 5'. Clean below 5'.

rigure		Kerinedy/Jenks/Chillon			
	na Industrial Proper	DATE 11/12/87 BORING B-2			
<u> </u>	38.00	_{BY} OL			
Details	of Boring and Well Cons	struct	ion		SURFACE ELEVATION:
SAMPLE NUMBER SAMPLER TYPE BLOWS/FINTERVAL	Well Construction	DEPTH IN FEET	nscs	SYMBOLS	DRILLING METHODS: P-50 truck-mounted, hollow stem auger. SAMPLING METHODS: Split barrel sampler. DRILLER: Rick Seiler (F.L.D.)
1 DRIVE 12		-1 -2 -3 -4 -5 -6	5	6	9 0' to 5' Sandy gravel fill, stained dark 0' to approx. 3'. Clean below 3'.

Figure

Kennedy/Jenks/Chilton

Figure			Refinedy/Jenks/Chillon		
PROJECT Tacom	a Industrial Proper	ties	5		DATE 11/12/87 BORING B-3
JOB NO 88673	8.00				BY OL BOTTO
Details o	f Boring and Well Cons	struc	tion	1	SURFACE ELEVATION:
SAMPLE NUMBER SAMPLER TYPE BLOWS/FT.INTERVAL	Well Construction		USCS	SYMBOLS	DRILLING METHODS: P-50 truck-mounted, hollow stem auger. SAMPLING METHODS: Split barrel sampler. DRILLER: Rick Seiler (F.L.D.)
		1 - 3 - 4 - 5 - 6			0 0' to 6' Sandy gravel fill, stained dark 0' to approx. 4'. Clean below 4'.

F 37

SHEET _____ OF ____

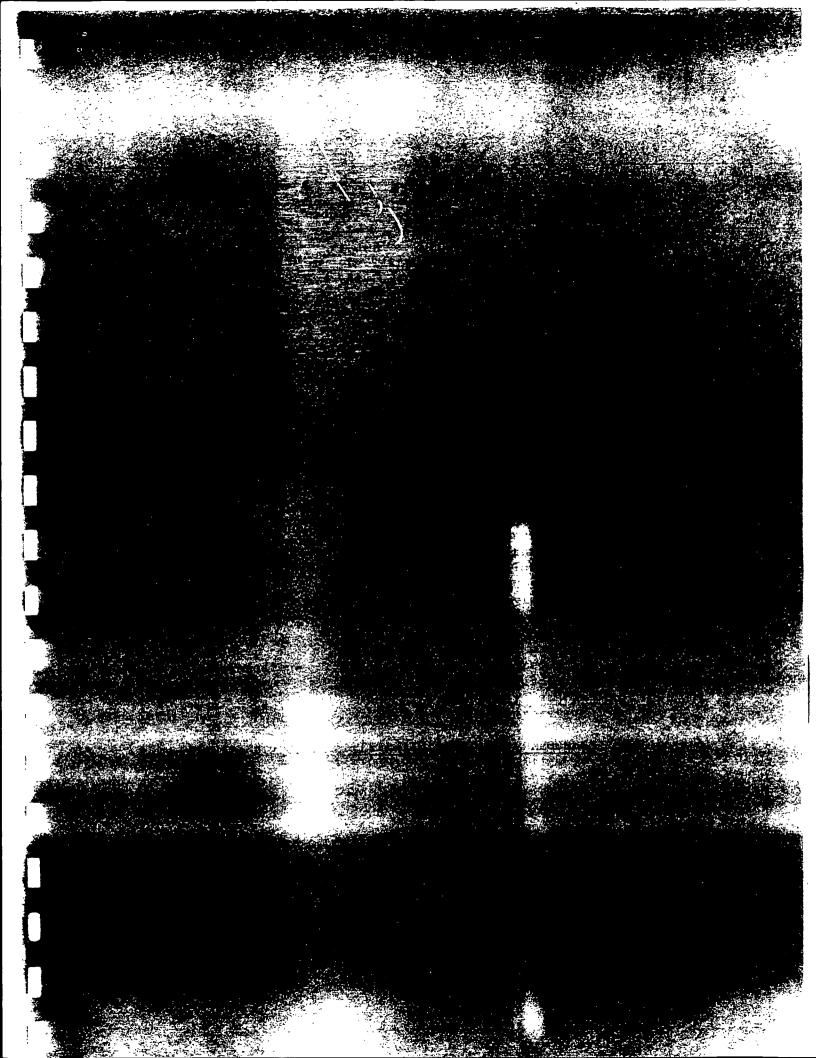
Figure

Figu	gure						Vermen's	/ Jenks/ Chillon	
PROJE	racoma industrial rioperties						DATE 11/12/87	BORING B-4	
JOP N	<u>o 88</u>	36738	3.00				ev OL	20,	
	Details of Boring and Well Construction						SURFACE ELEVATION:		
EA	≝						P-50 truck-mounted hollow stem auger.		
SAMPLE NUWRER	SAMPLER TYPE	BLOWS/FT:INTERVAL		DEPTH IN FEET		Ŋ		Split barrel sampler.	
MPLE	MPLE	OWS/F		H	SOSO	SYMBOLS			
\$	\$	9	1111	-	1 3	<u> </u>	DRILLER:	Rick Seiler (F.L.D.)	
i				-1 -2				A.C. Pavement.	
				-3			@ 2" to 10½"	Sand, fine brown to grey, moist, gravel < 5%.	
				-4				Boring terminated at 10½'	
1	DRIVE	7,10 & 13 for 5" each		-5 -6	SM			No product odor or unusual color in samples or cuttings.	
		p eacr		-7 -8				Probed adjacent fuel tank bottom @ 8'; tank contains	
1				-9				approx. 3 3/4" liquid. Odor-gasoline.	
	-	10.13		-10		: : :			
2	DRIVE	for 6"	1 11	-11					
		each		-12					
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POJE	o Ta	Coma	Industr	ial Propert	ies			DATE 11/2/87	BORING MW-5
JOP NO 886738.00						BY OL	BORING PW 3		
ı	Deta	ils of	Boring a	nd Well Cons	struc	tior	1	SURFACE ELEVATION	· · · · · · · · · · · · · · · · · · ·
SAMPLE NUMBER	SAMPLER TYPE	BLOWS/FT:INTERVAL	Tradfic	enstruction rated monument right locking cap.	DEPTH IN FEET		N.S		P-50 truck-mounted, hollow stem auger. S: Split barrel sampler.
SAMPL	SAMPL	BLOWS	*		06 <i>PT</i> H	uscs	SYMBOLS	DRILLER:	Rick Seiler (F.L.D.)
					-1			0 0" to 2"	A.C. Pavement.
		7.13 &		Backfilled with drill cuttings	-2 -3 -4	SW		9 2" to 3'	Gravelly sand, fill, debris, slag. Gravels 85%, sand 15%. dark brown - black. gravels up to 2" diam.
1	DRIVE	15 for 6" each		Bentonite	-6 -7			₽ 5' to 6'⅓	Gravelly sand, well graded, moist, Sand 60%, Gravel 40%.
				seal. Approx. 7' to 15½'.	-8 -9 -10	SM		0 10' to 11⅓'	Gravel and sand, Sand 50%, Gravel 50%. Gravels < 1" diam, rounded. Sand brown- tan.
2	DRIVE	10,14 E 20 for 6" each			-11 -12 -13				
3	DRIVE	7,15 & 16 for 6" each		Top of sand @ 15½'.	-15 -16 -17 -18	SW		@ 15' to 16½'	Gravelly sand, Sand 90%, Gravel 10%. Gravels < 1" diam; sand fine to medium, angular.
4	DRIV	16.21 E& 42 _for 6"		Top of screen @ 19½'.	-19 -20 -21		00,	0 20' to 21½'	Gravelly sand, Coarse sand 65%, Gravel 35%, gravels < 낮 diam, moist.
		each		10' to 20' Filter sand. 2" diam PVC	-22 -23		ە ي		Gravelly sand, Coarse sand 65%, Gravel 35%, gravels < 낮 diam, moist.
_		32,35		screen with 0.02" slot.	-24 -25 -26	SW	995	v @ 30' to 31½'	Gravelly sand, Sand 50%, Gravel 50%.
5	DRIV	E& 30 _for 6" each			-27 -28			0	No unusual odor or color in samples or drilling cuttings below 5'.
					-29 -30		,0		Boring terminated at 40'.
6	DRIV	48 for E 6" 50 for 5"			-31 -32		6°		

Figure

Details of Boring and Well Construction Well Construction Boring MW-5 Well Construction Boring MW-5 Bori		PROJECT Tables Tools at 12 D				recinica y				
Details of Boring and Well Construction Well Construction Surface Elevation: DRILLING METHODS: SAMPLING METHODS: SAM		006739 00						OI .	BORING	MW-5
Well Construction Solution S		`			struc	ction	1	, , , , , , , , , , , , , , , , , , ,		
33 -34 -35 -36 -36 -37 -39 -40								1		
33 -34 -35 -36 -36 -37 -39 -40	E NUMB	ER TYPI	/FT-INTE		IN FEET		S	SAMPLING METHODS:		
Bottom of screen 0 37 39½ -38 End cap39 -40	SAMPL	SAMPL	BLOWS		ОЕРТН	USCS	SYMBO	DRILLER:	Rick Seiler (F.L	.D.)
				screen @ 39½' End cap.	-33 -34 -35 -36 -37 -38 -39				Rick Seiler (F.L	.0.)



APPENDIX B

CALCULATION OF ESTIMATED PERMISSIBLE
CONCENTRATION AND MAXIMUM ACCEPTABLE WATER CONCENTRATIONS
FOR COMPOUNDS AT FORMER IRON FOUNDRY SITE

EPCs: EPA has not established a multimedia environmental goal for $\overline{1,1,2}$ -Trichloro-1,2,2-trifluoroethane. The development of the suggested action level for this compounds in soil and groundwater presented below follows EPA formulas for calculating estimated permissible concentrations (EPC)^a:

Health Effects in Water (WH2)C

WH2(ug/L) = $13.8 \times TLV(mg/m^3)$

Health Effects on Land (LH)C

LH(mq/Kq) = 200 WH2(mg/L)

Example:

The Threshold Limit Value (TLV) for 1,1,2-Trichloro-1,2,2-trifluoroethane is $7600~\text{mg/m}^3$ b. The calculation of WH2 and LH is therefore:

WH2 = $13.8 \times 7600 = 104,880 \text{ ug/l}$

LH = 200 x 104,880 x $\frac{1}{1000}$ = 20,976 mg/Kg

Our suggested soil action level is set at 20,976 mg/Kg. Our suggested water action level is set at 104,880 mg/l.

Notes:

- a EPA, 1980. "Multimedia Environmental Goals for Environmental Assessment," Volume I (Supplement A). EPA 600/7-80-041.
- b TLVs, Threshold Limit Values for Chemical Substances and Physical Agents in the Work Environment with Intended Changes for 1986-87. American Conference of Governmental Industrial Hygienists, 1986.
- C WH2 Health Effects in Water

LH - Health Effects on Land

TLV - Threshold Limit Value

Nomenclature used by EPA in note a

MAWCs: EPA has developed a methodology for calculating maximum acceptable concentrations in a media for carcinogenic or suspected carcinogenic compounds. These concentrations are based on the potency slope developed by EPA's Carcinogen Assessment Group (CAG) and the selected risk level. EPA typically uses risks on the order of 10⁻⁴ to 10⁻⁷ when selecting an acceptable risk level.

To calculate the maximum acceptable water concentrations (MAWC), the acceptable dose (AD) must first be calculated using the following equation:

AD (mg/kg/day) = Risk Level/Potency Slope

To assess the AD for a typical 70 kg adult, the AD was multiplied by 70 kg, resulting in an AD in mg/day. The MAWC was then calculated by the following equation:

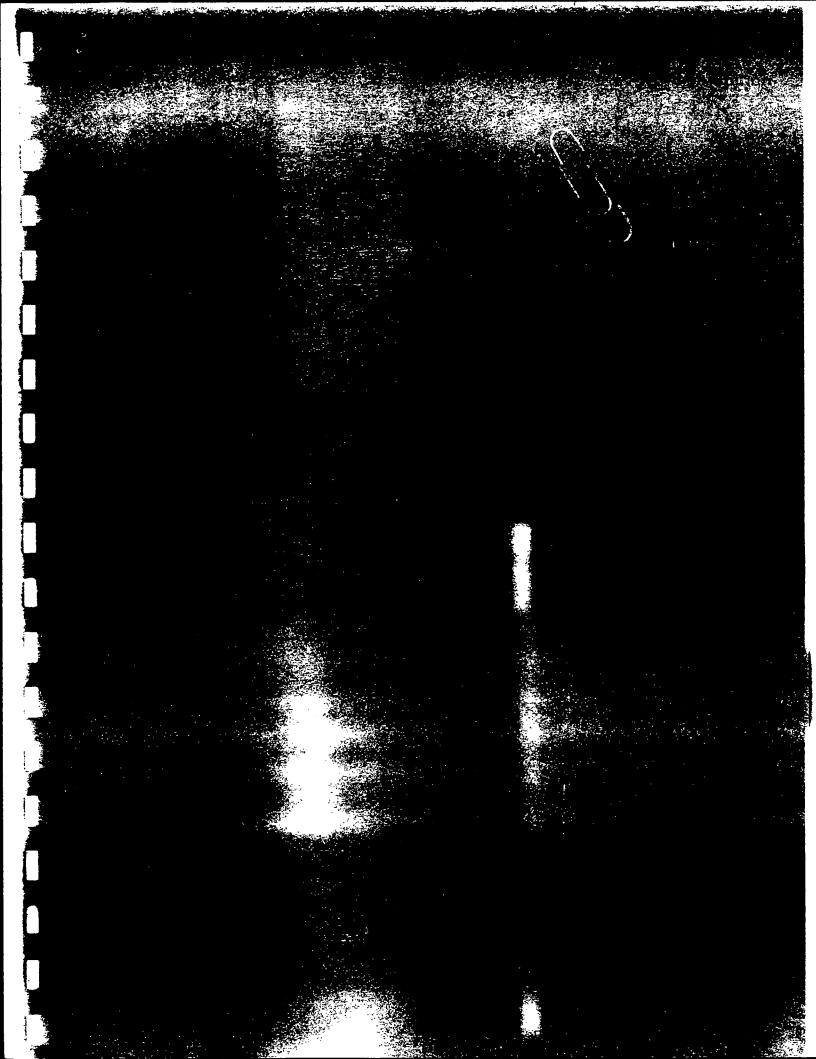
MAWC (mg/1) = AD (mg/day)/Water Ingestion Rate (1/day)

EPA has used two liters per day as the average daily water intake of a human adult.a

MAWCs were calculated for tetrachloroethylene. The potency slope for tetrachloroethylene is 5.1×10^{-2} mg/kg/day.^b Calculating the MAWC for a risk level of 10^{-5} led to a concentration of 0.07 mg/l for tetrachloroethylene.

Notes:

- a EPA, 1985. "EPA Proposal to Set Recommended Drinking Water Limits for Inorganic, Synthetic Organic Chemicals, Microorganisms." 50 FR 46936, November 13, 1985.
- b EPA, 1985. Office of Health and Environmental Assessment. "Health Assessment Document for Tetrachloroethylene (Perchloroethylene)," Washington, D.C.



Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For Attention Kennedy/Jenks/Chilton

Mr. Nathan Graves

33301 Ninth Avenue South, Suite 100 Address Federal Way, WA 98003

Received 10/7/86 Reported 11/3/86 (K/J/C 6738)

866000 Lab. No.

Source TIP

Tacoma, WA Date Collected

Time Collected

Collected by

Sample Type:

Soil, GS-2

10/2/86

K/J/C

Soil, GS-2

PRIORITY POLLUTANT COMPOUNDS

Volatiles	ug/kg(ppb)	Volatiles	ug/kg(ppb)
chloromethane	<100	1,2-dichloropropane	<100
bromomethane	<100	trans-1,3-dichloropropylene	<100
vinyl chloride	<100	trichloroethylene	<100 ·
chloroethane	<100	benzene	<100
methylene chloride	(1)	dibromochloromethane	<100
trichlorofluoromethane	<100	cis-1,3-dichloropropylene	<100
1.1-dichloroethylene	<100	1,1,2-trichloroethane	<100
1,1-dichloroethane	<100	2-chloroethylvinyl ether	<100
1,2-dichloroethylene	<100	bromoform	<100
chloroform	(1)	tetrachloroethylene	300
	<100	1,1,2,2-tetrachloroethane	<100
1,2-dichloroethane 1,1,1-trichloroethane	550	toluene	<100
carbon tetrachloride	<100	chlorobenzene	<100
bromodichloromethane	<100	ethylbenzene	<100
	NON-PRIORITY P	OLLUTANT COMPOUNDS	
acetone	<1000	vinyl acetate	<100
carbon disulfide	<100	4-methyl-2-pentanone	<1000
1,1,2-Trichloro-		2-hexanone	<1000
1 2 2-trifluoroethane	<100	styrene	<100

xylenes

Comments: (1) This component was not quantifiable in this sample.

<2000

Analysis by U.S. EPA Method 8240, reported in micrograms per kilogram, wet (as received) weight basis.

Analyst EW, BK

1,2,2-trifluoroethane

methyl ethyl ketone

5000

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For

Kennedy/Jenks/Chilton

Attention

Mr. Nathan Graves

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 10/7/86 Reported 12/3/86 (K/J/C 6738)

Lab.No. 866000

Date Collected 10/2/86

Collected by K/J/C.

Source TIP, Tacoma, WA

Time Collected -

Sample Type: GS-2

		POLLUTANT COMPOUNDS	10
COMPOUNDS	ug/kg(ppl		ug/kg(ppb
phenol	<200	2,6-dinitrotoluene	<200
2-chlorophenol	<200	acenaphthene	<200
2-nitrophenol	<200	2,4-dinitrotoluene	<200
2,4-dimethylphenol	<200	fluorene	<200
2,4-dichlorophenol	<200	diethyl phthalate	<200
4-chloro-3-methylphenol	<200	4-chlorophenylphenyl ether	<200
2,4,6-trichlorophenol	<200	N-nitrosodiphenylamine	<200
2,4-dinitrophenol	<1000	1,2-diphenylhydrazine (azobenzene)	<200
2-methyl-4,6-dinitrophenol	<1000	4-bromophenylphenyl ether	<200
4-nitrophenol	<200	hexachlorobenzene	<200
pentachlorophenol	<200	phenanthrene	<200
bis(2-chloroethyl)ether	<200	anthracene	<200
1,3-dichlorobenzene	<200	di-n-butyl phthalate	<200
1,4-dichlorobenzene	<200	fluoranthene	<200
1,2-dichlorobenzene	<200	benzidine	<1000
bis(2-chloroisopropyl)ether	<200	pyrene	<200
hexachloroethane	<200	benzyl butyl phthalate	<200
N-nitrosodi-n-propylamine	<200	benz[a]anthracene	<200
nitrobenzene	<200	3,3'-dichlorobenzidine	<400
isophorone	<200	chrysene	<200
bis(2-chloroethoxy) methane	<200	bis(2-ethylhexyl)phthalate	<200
1,2,4-trichlorobenzene	<200	di-n-octyl phthalate	<200
naphthalene	<200	benzo[b]fluoranthene	<200
hexachlorobutadiene	<200	benzo[k]fluoranthene	<200
hexachlorocyclopentadiene	<200	benzo[a]pyrene	<200
2-chloronaphthalene	<200	indeno[1,2,3-cd]pyrene	<200
dimethyl phthalate	<200	dibenz[a,h]anthracene	<200
acenaphthylene	<200	benzo[ghi]perylene	<200
	NON-PRIORIT	Y POLLUTANT COMPOUNDS	
benzoic acid	<1000	4-chloroaniline	<1000
2-methylphenol	<200	2-methylnaphthalene	<200
4-methylphenol	<200	2-nitroaniline	<1000
2,4,5-trichlorophenol	<200	3-nitroaniline	<1000
aniline	<200	dibenzofuran	<200
benzyl alcohol	<400	4-nitroaniline	<1000

Analysis by U.S. EPA Method 8270, reported in micrograms per kilogram, wet (as received) weight basis.

Note: Extract was oily, requiring dilution before analysis

Analyst BK

Manager Zeveel

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

GC/MS Volatiles Surrogate Standard Recovery Report

For Attention

Kennedy/Jenks/Chilton

Attention Address

Mr. Nathan Graves 33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/9/86 11/3/86

(K/J/C 6738)

Sample Ide	entification	Percent R			
Lab No.	Type	1,2-Dichloroethane-d4	Fluorobenzene	4-Bromofluorobenzene	
865999	Soil	104	78	124	
866000	Soil	109	78	51	

Analyst BK Manager Levell R. Smith

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Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For

Kennedy/Jenks/Chilton

Attention Mr. Nathan Graves

Address 33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 10/7/86 Reported 12/3/86

(K/J/C 6738)

Lab.No. 866001

Date Collected 10/2/86

Collected by K/J/C.

Source TIP, Tacoma, WA

Time Collected 1241

Sample Type: SS-4

	PRIORITY PO	LLUTANT COMPOUNDS	
COMPOUNDS	ug/kg(ppb)	COMPOUNDS	ug/kg(ppb)
phenol	<2000	2,6-dinitrotoluene	<2000
2-chlorophenol	<2000	acenaphthene	<2000
2-nitrophenol	<2000	2,4-dinitrotoluene	<2000
2,4-dimethylphenol	<2000	fluorene	<2000
2,4-dichlorophenol	<2000	diethyl phthalate	<2000
4-chloro-3-methylphenol	<2000	4-chlorophenylphenyl ether	<2000
2,4,6-trichlorophenol	<2000	N-nitrosodiphenylamine	<2000
2,4-dinitrophenol	<10,000	1,2-diphenylhydrazine (azobenzene)	<2000
2-methyl-4,6-dinitrophenol	<10,000	4-bromophenylphenyl ether	<2000
4-nitrophenol	<2000	hexachlorobenzene	<2000
pentachlorophenol	<2000	phenanthrene	<2000
bis(2-chloroethyl)ether	<2000	anthracene	<2000
1,3-dichlorobenzene	<2000	di-n-butyl phthalate	<2000
1.4-dichlorobenzene	<2000	fluoranthene	₹2000
1,2-dichlorobenzene	<2000	benzidine	<10,000
bis(2-chloroisopropyl)ether	<2000	pyrene	<2000
hexachlorosthane	<2000	benzyl butyl phthalate	<2000
N-nitrosodi-n-propylamine	<2000	benz[a]anthracene	<2000
nitrobenzene	<2000	3,3'-dichlorobenzidine	<4000
isophorone	<2000	chrysene	<2000
bis(2-chloroethoxy) methane	<2000	bis(2-ethylhexyl)phthalate	<2000
1,2,4-trichlorobenzene	<2000	di-n-octyl phthalate	<2000
naphthalene	<2000	benzo[b]fluoranthene	<2000
hexachlorobutadiene	<2000	benzo(k)fluoranthene	<2000
hexachlorocyclopentadiene	<2000	benzo[a]pyrene	<2000
2-chloronaphthalene	<2000	indeno[1,2,3-cd]pyrene	<2000
dimethyl phthalate	<2000	dibenz[a,h]anthracene	<2000
acenaphthylene	<2000	benzo[ghi]perylene	<2000
accinapitatily 20110			
	NON-PRIORITY	POLLUTANT COMPOUNDS	410,000
benzoic acid	<10,000	4-chloroaniline	<10,000
2-methylphenol	<2000	2-methylnaphthalene	<2000
4-methylphenol	<2000	2-nitroaniline	<10,000
2,4,5-trichlorophenol	<2000	3-nitroaniline	<10,000
aniline	<2000	dibenzofuran	<2000
benzyl alcohol	<4000	4-nitroaniline	<10,000

Analysis by U.S. EPA Method 8270, reported in micrograms per kilogram, wet (as received) weight basis.

Note: Extract was oily, requiring dilution before analysis.

Analyst BK

anager weell

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415 362 6065

For

Kennedy/Jenks/Chilton

Received

10/23/86

Attention

Nathan Graves

Reported

12/3/86

Address

33301 Ninth Avenue South, Suite 100 Federal Way, WA 98003

eported

(K/J/C 6738)

Lab. No.

867367

Source

Soil I.D.: SS-6

TIP

Tacoma, WA

Date Collected

10/17/86

Time Collected

1450

Collected by

K/J/C

Analysis (1)

Units

Analytical Results

Polychlorinated

biphenyls (PCBs)

mg/Kg

<1

Comments: (1) Analysis by EPA Method 8080; reported in milligrams per kilogram, wet (as received) weight basis. Extraction by EPA Method 3540 (Soxhlet). Reported as Aroclor 3540.

Reference: "Test Methods for Evaluating Solid Waste — Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and "California Administrative Code Title 22, Div. 4".

Analyst _____

MC

Manager

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The hability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention Kennedy/Jenks/Chilton

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/23/86 10/31/86

(K/J/C 6733)

Lab. No.

867370

Source TIP Soil I.D.: SD-1

Tacoma, WA

Date Collected

10/9/86

Time Collected

1608

Collected by

K/J/C

Analysis

Units

Analytical Results

Lead (Pb) (2)

mg/Kg (1) 320

(1) Milligrams per kilogram, wet (as received) weight basis. (2) Analysis by atomic absorption spectrophotometer.

Reference: "Test Methods for Evaluating Solid Waste - Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and "California Administrative Code Title 22, Div. 4".

Analyst .

Manager

Nathan Graves, K/J/C, Federal Way, WA

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Water Analysis Report

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For

Kennedy/Jenks/Chilton

Attention Owen Loshbough

33301 Winth Avenue South, Suite 100 Address

Federal Way, WA 98003

Received Reported

Analytical Results

10/23/86 10/31/86

(K/J/C 6733)

Lab. No.

867371

Source

Water I.D.: SD-2

TIP

Tacoma, WA

Date Collected

10/9/86

Time Collected

1616

Collected by

K/J/C

Units Analysis mg/L 9.0 Lead (Pb)

Comments:

Analysis by: "Standard Methods for the Examination of Water and Wastewater", Current Edition APHA

Analyst

Manager .

Nathan Graves, K/J/C, Federal Way, WA This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For

Kennedy/Jenks/Chilton

Received

10/23/86

Attention

Owen Loshbough

Reported

10/31/86

Address

33301 Ninth Avenue South, Suite 100 Federal Way, WA 98003

(K/J/C 6733)

Lab. No.

867368

Source

Soil I.D.: SD-3

TIP

Tacoma, WA

Date Collected

10/10/86

Time Collected

1027

Collected by

K/J/C

Analysis

Units

Analytical Results

Lead (Pb) (2)

mg/Kg (1) 530

Comments: (1) Milligrams per kilogram, wet (as received) weight basis. (2) Analysis by atomic absorption spectrophotometer.

Reference: "Test Methods for Evaluating Solid Waste -- Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and 1984 "California Administrative Code Title 22, Div. 4".

AD Analyst

Manager

cc: Nathan Graves, K/J/C, Federal Way, WA

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Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention

Kennedy/Jenks/Chilton

Nathan Graves

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/23/86

12/3/86

(K/J/C 6738)

Lab. No.

867368

Source.

Soil I.D.:

SD-3

TIP

Tacoma, WA

Date Collected

10/10/86

Time Collected:

1021

Collected by

K/J/C

Analysis-

Units

Analytical Results:

Petroleum Hydrocarbons

(as diesel fuel)

mg/Kg

8.5

Comments: Analysis of pentane extract by gas chromatography with flame ionization detection, using commercial hydrocarbon samples as comparison standards. Results reported in milligrams per kilogram, wet (as received) weight basis.

Reference: "Test Methods for Evaluating Solid Waste — Physical/Chemical Methods", SW—846, Second Edition (Revised 1984), and: "California Administrative Code Title 22, Div. 4".

Analyst ____ KA

Manager

Leverett R. Smith

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Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention Kennedy/Jenks/Chilton

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

10/23/86 Received Reported

10/31/86

(K/J/C 6733)

Lab. No.

867372

Source · TIP Soil I.D.:

SD-4

Tacoma, WA

Date Collected

10/9/86

Time Collected

1633

Collected by

K/J/C

Analysis

Units

Analytical Results

Lead (Pb) (2)

mg/Kg (1)

Comments: (1) Milligrams per kilogram, wet (as received) weight basis is (2) Analysis by atomic absorption spectrophotometer. Reference: "Test Methods for Evaluating Solid Waste - Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and "California Administrative Code Title 22, Div. 4". Analyst _ Manager

cc: Nathan Graves, K/J/C, Federal Way, WA
This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof

Water Analysis Report

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention

Address

Kennedy/Jenks/Chilton

Owen Loshbough

Owen Losnbough

33301 Ninth Avenue South, Suite 100 Federal Way, WA 98003

Received Reported 10/23/86 10/31/86

(K/J/C 6733)

Lab. No.

867373

Source

Water I.D.:

SD-5

TIP Tacoma, WA

Date Collected

10/9/86

Time Collected

640

Collected by

K/J/C

Analysis Units Analytical Results

Lead (Pb) mg/L 1.6

er salete

er 😘

Comments:

11/ June 1

Analysis by: "Standard Methods for the Examination of Water and Wastewater", Current Edition, APHA

AD

"Manage

Leverett R. Smith

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Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention

Kennedy/Jenks/Chilton

on Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/23/86 10/31/86

(K/J/C 6733)

Lab. No.

867374

Source

Soil I.D.:

SD-6

TIP

Tacoma, WA

Date Collected

10/9/86

Time Collected

1651

Collected by

K/J/C

Analysis Units

Lead (Pb) (2) mg/Kg (1)

1200

in the second converses

Analytical Results

A STATE OF THE STA

ordong

Comments: (1) Milligrams per kilogram, wet (as received) weight basis. (2) Analysis by atomic absorption spectrophotometer.

Reference: "Test Methods for Evaluating Solid Waste — Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and "California Administrative Code Title 22, Div. 4".

Analyst ____AD

Manager

cc: Nathan Graves, K/J/C, Federal Way, WA
This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples.
The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons

so informed of the contents hereof.

EP Toxicity Analysis Report

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention Address Kennedy/Jenks/Chilton

Mr. Nathan Graves

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/23/86 12/3/86

(K/J/C 6738)

Lab. No.

867374

Source

SD-6

TIP

Tacoma, WA

Date Collected

10/9/86

Time Collected

1651

Collected by

K/J/C

Contaminant

Concentration in Extract (1)

Lead (Pb)

mg/L

Units

<0.5

Final pH of extract = 5.2

Comments:

Reference: "Test Methods for Evaluating Solid Waste - Physical/Chemical Methods,"

SW-846, Second Edition, 1984, U.S. EPA.

Analyst TK, AI

Manager .

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Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention Kennedy/Jenks/Chilton

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received Reported 10/23/86 10/31/86

(K/J/C 6733)

Lab. No.

867369

Source TIP Soil I.D.:

SD-7

Tacoma, WA

Date Collected

10/9/86

Time Collected

1701

Collected by

K/J/C

Analysis

Units

Analytical Results

netwrical Results

mg/Kg(1)Lead (Pb) (2) and (-310

sa annan gan a

(1) Milligrams per kilogram, wet (as received) weight basis.

(2) Analysis by atomic absorption spectrophotometer bushs-

Reference: "Test Methods for Evaluating Solid Waste - Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and "California Administrative Code Title 22, Div. 4".

AD Analyst

Manager

Nathan Graves, K/J/C, Federal Way, WA This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons

so informed of the contents hereof,

and the fall that

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

10/12/06

Attention Address	Nathan Graves 33301 Ninth Avenue S Federal Way, WA 980	outh, Suite 100	Reported	12/3/86 12/3/86 (K/J/C 6738)
Lab. No.	redefal way, wa 500	867369		(2,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0
Source TIP Tacoma,	Soil I.D.:	SD-7		
Date Collect	ted	10/9/86	Calums	n
Time Collec	ted	1701	n is some	
Collected by		V / 1 / C		•

Analysis Units Analytical Results

Petroleum Hydrocarbons
(kerosene) mg/Kg 61

Comments: Analysis of pentane extract by gas chromatography with flame ionization detection, using commercial hydrocarbon samples as comparison standards. Results reported in milligrams per kilogram, wet (as received) weight basis.

Reference: "Test Methods for Evaluating Solid Waste - Physical/Chemical Methods", SW-846, Second Edition (Revised 1984), and

"California Administrative Code Title 22, Div. 4".

Analyst KA Manager Tevel K. Smith

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For

Kennedy/Jenks/Chilton

Attention Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 11/17/87

Reported 11/30/87 (K/J/C 866738.00)

Lab. No. 877719

Source

TIP

Date Collected

11/12/87

Time Collected

1352

Date Analyzed: 11/24/87

Collected by

K/J/C

Sample Type:

Soil: B-2, Depth 5-6.5 ft 200 8

PRIORITY POLLUTANT COMPOUNDS

•			a rate material in Various and		
Volatiles	ug/Kg(Volatiles: # Land	ug/Kg	g(ppb)
. ;		t. Lin	An all killing all arrivers		Det. Lim.
chloromethane	<10	10	bromodichloromethane Trovaled a	<5	5 ·
bromomethane	<10	10	1,2-dichloropropane	<5	5
vinyl chloride	<10	10	trans-1,3-dichloropropylene	<5	5
chloroethane	<10	10	trichloroethylene	<5	5
methylene chloride	<20	20	benzene - opropyteme	<5	5
acrolein -	<30	30	dibromochloromethane	<5	5 5
acrylonitrileth.	<10	10	cis-1,3-dichloropropylene	<5	5
trichlorofluctomethane	<5	5	1,1,2-trichloroethane	<5	5
1,1-dichloroethylene	<5	5	2-chloroethylvinyl ether	<5	5
1,1-dichloroethane	<5	5	bromoform	<5	5
1,2-dichloroethylene	<5	5	tetrachloroethylene	<5	5
Chloroform	<5	5	1,1,2,2-tetrachloroethane	<5	5
1,2-dichloroethane	<5	5	toluene	<5∙	5
1,1,1-trichloroethane	<5	5	chlorobenzene	<5	5
carbon tetrachloride	<5	5	ethylbenzene	<5	5
	NON-PRIO	RITY PO	DLLUTANT COMPOUNDS		
acetonitrile	<30	30	vinyl acetate	<10	. 10
acetone	<10	10	4-methyl-2-pentanone	<10	- 10
carbon disulfide	<5	5	2-hexanone	<10	10
1,1,2-Trichloro-			styrene	<5	5
1,2,2-trifluoroethane	7	5	xylenes	<5	5
methyl ethyl ketone	<10	10	And the second of the second o		
- · · · · · · · · · · · · · · · · · · ·			•		

Comments: Analysis by U.S. EPA Method 8240, reported in micrograms per kilogram, wet (as received) weight basis.

Analyst TJL, SL

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For

Kennedy/Jenks/Chilton

Attention

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received

Reported 11/30/87

(K/J/C 866738.00)

Lab. No. Method Blank

Source

Date Collected

Time Collected

Date Analyzed: 11/24/87

Collected by

K/J/C

Sample Type:

Water:

PRIORITY POLLUTANT COMPOUNDS

	PRIURI	II PUL	CUIANI COHPOUNDS	•	
<u>Volatiles</u>	ug/L (ppb) Volatiles		ug/L (ppb)		
	De	et. Lin	n.		et. Lim.
chloromethane	<10	10	bromodichloromethane	<5	5
bromomethane	<10	10	1,2-dichloropropane	<5	5
vinyl chloride	<10	10	trans-1,3-dichloropropylene	<5	5
chloroethane	<10	10	trichloroethylene	<5	5
methylene chloride	7	5	benzene	< 5	5
acrolein	<30	30	dibromochloromethane	<5	5
acrylonitrile	<10	10	cis-1,3-dichloropropylene	<5	5
trichlorofluoromethane	<5	5	1,1,2-trichloroethane	<5	5
1,1-dichloroethylene	<5	5	2-chloroethylvinyl ether	<5	5
l,l-dichloroethane	<5	5	bromoform	<5	5
l,2-dichloroethylene	<5	5	tetrachloroethylene	<5	5
Chloroform	<5	5	1,1,2,2-tetrachloroethane	<5	5
1,2-dichloroethane	<5	5	toluene	<5	5
1,1,1-trichloroethane	<5	5	chlorobenzene	<5	5
carbon tetrachloride	<5	. 5	ethylbenzene	<5	5
	NON-PRIO	RITY PO	OLLUTANT COMPOUNDS		
acetonitrile	<30	30	vinyl acetate	<10	10
acetone	<10	10	4-methyl-2-pentanone	<10	10
carbon disulfide	<5	5	2-hexanone	<10	10
1,1,2-Trichloro-			styrene	<5	5
1,2,2-trifluoroethane	<5	5	xylenes	<5	5
methyl ethyl ketone.	<10	10	÷.		
· •				-	

Comments: Analysis by U.S. EPA Method 8240, reported in micrograms per liter.

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

GC/MS Volatiles Surrogate Standard Recovery Report

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

For Attention Kennedy/Jenks/Chilton

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 11/17/87 Reported 11/30/87

(K/J/C 866738.00)

Sample Identi	fication	Pe:	rcent Recoveri	es
Lab No.	Type	1,2-Dichloroethane-d4	Toluene-d8	4-Bromofluorobenzene
877719	Soil	98	107	87
877719 Dup.	Soil	102	103	88

Acceptable Recoveries:	<u>Water</u>	<u>Soil</u>
1,2-Dichloroethane-d4	76-114	70-121
Toluene-d8	88-110	81-117
4-Bromofluorobenzene	86-115	74-121

TJL, SL

Manager

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

Soil Analysis Report

For:

Kennedy/Jenks/Chilton

Attn:

Owen Loshbough

Address:

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

(K/J/C 866738.00)

Received 11/17/87

Reported 11/30/87

Lab. No.:

877720

Source:

Sample I.D.:

TIP

Depth: 9-10.5 ft

B-4

Tacoma, WA

Date Collected:

11/12/87

Date Analyzed: 11/24/87

Time Collected:

1445

Collected by:

K/J/C

Units*		Analytical Results	Det. Limit
	-		
ug/Kg	<1		1.
ug/Kg	<1		1
ug/Kg	<1		ī
ug/Kg	<1	•	ī
ug/Kg	<1	·	ī
ug/Kg	<1		1
ug/Kg	<1	· ·	$\bar{1}$
ug/Kg	<1		ī
ug/Kg	<1		$\overline{1}$
ug/Kg	<1		1
	ug/Kg ug/Kg ug/Kg ug/Kg ug/Kg ug/Kg ug/Kg ug/Kg	ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1	ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1 ug/Kg <1

Comments: Analysis by EPA Method 8020 (Purgeable Aromatics)

* Micrograms per kilogram, wet (as received) weight basis.

Analyst DC, SL

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

94105 San Francisco, CA

415-362-6065

For

Kennedy/Jenks/Chilton

Attention

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 11/18/87 Reported 11/25/87

(K/J/C 866738.00)

Lab. No. 877736

Source

TIP

Date Collected

11/16/87

Time Collected

1530

Date Analyzed: 11/20/87

Collected by

K/J/C

Sample Type:

Water: MW-5

PRIORITY POLLUTANT COMPOUNDS

Volatiles	ug/L	(ppb)	Volatiles	ug/L	(ppb)
	Det. Lim.		1.		et. Lim.
chloromethane	<20	20	bromodichloromethane	<10	10
bromomethane	<20	20	1,2-dichloropropane	<10	10
vinyl chloride	<20	20	trans-1,3-dichloropropylene	<10	10
chloroethane	<20	20	trichloroethylene	<10	10
methylene chloride	<15	15	benzene	<10	10
acrolein	<60	60	dibromochloromethane	<10	10
acrylonitrile	<20	20	cis-1,3-dichloropropylene	<10	10
trichlorofluoromethane	200	10	1,1,2-trichloroethane	<10	10
l,l-dichloroethylene	<10	10	2-chloroethylvinyl ether	<10	10
1,1-dichloroethane	<10	10	bromoform	<10	10
1,2-dichloroethylene	<10	10	tetrachloroethylene	<10	10
Chloroform	<10	10	1,1,2,2-tetrachloroethane	<10	10
1,2-dichloroethane	<10	10	toluene	<10	10
1,1,1-trichloroethane	34	10	chlorobenzene	<10	10
carbon tetrachloride	<10	10	ethylbenzene	<10	10
	NON-PRIO	RITY PO	DLLUTANT COMPOUNDS		
acetonitrile	<60	60	vinyl acetate	<20	20
acetone	<20	20	4-methyl-2-pentanone	<20	20
carbon disulfide	<10	10	2-hexanone	<20	20
1,1,2-Trichloro-			styrene	<10	10
1,2,2-trifluoroethane	<10	10	xylenes	<10	. 10
methyl ethyl ketone	<20	20			

Analysis by U.S. EPA Method 8240, reported in micrograms per liter. Comments:

Analyst TJL, SL

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton, Laboratory Division

657 Howard Street

San Francisco, CA 94105

415-362-6065

For

Kennedy/Jenks/Chilton

Attention

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received

Reported 11/25/87

(K/J/C 866738.00)

Lab. No.

Method Blank

Source

Date Collected

Time Collected

Date Analyzed: 11/19/87

Collected by

K/J/C

Sample Type:

Water:

PRIORITY POLLUTANT COMPOUNDS

Volatiles	ug/L ((ppb)	Volatiles	ug/L	(ppb)
	D€	et. Li	.m.		Det. Lim.
chloromethane	<10	10	bromodichloromethane	<5	5
bromomethane	<10	10	1,2-dichloropropane	<5	5
vinyl chloride	<10	10	trans-1,3-dichloropropylene	< 5	5
chloroethane	<10	10	trichloroethylene	<5	5
methylene chloride	<5	5	benzene	<5	5
acrolein	<30	30	dibromochloromethane	< 5	5
acrylonitrile	<10	10	cis-1,3-dichloropropylene	<5	5
trichlorofluoromethane	<5	5	1,1,2-trichloroethane	<5	5
1,1-dichloroethylene	<5	5	2-chloroethylvinyl ether	<5	5
1,1-dichloroeinane	<5	5	bromoform	<5	5
1,2-dichloroethylene	<5	5	tetrachloroethylene	<5	5
Chloroform	<5	5	1,1,2,2-tetrachloroethane	<5	5
1,2-dichloroethane	<5	5	toluene	<5	5
1,1,1-trichloroethane	. <5	5	chlorobenzene	<5	5
carbon tetrachloride	<5	5	ethylbenzene	<5	5
	NON-PRIO	RITY E	POLLUTANT COMPOUNDS		
acetonitrile	<30	30	vinyl acetate	<10	10
acetone	<10	10	4-methyl-2-pentanone	<10	10
carbon disulfide	<5	5	2-hexanone	<10	10
1,1,2-Trichloro-			styrene	<5	5 .
1,2,2-trifluoroethane	<5	5	xylenes	<5	5
methyl ethyl ketone	<10	10	•		

Comments: Analysis by U.S. EPA Method 8240, reported in micrograms per liter.

Analyst TJL, SL

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Kennedy/Jenks/Chilton

Laboratory Division

657 Howard Street San Francisco, California 94105 415-362-6065

GC/MS Volatiles Surrogate Standard Recovery Report

For Attention

Kennedy/Jenks/Chilton

Owen Loshbough

Address

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

Received 11/18/87 Reported 11/25/87

(K/J/C 866738.00)

Sample Ident	tification	Pe	rcent Recoveri	es ·
Lab No.	Type	1,2-Dichloroethane-d4	Toluene-d8	4-Bromofluorobenzene
877636	Water	104	97	97

Acceptable Recoveries:	Water	<u>Soil</u>	
1,2-Dichloroethane-d4	76-114	70-121	
Toluene-d8	88-110	81-117	
4-Bromofluorobenzene	86-115	74-121	

Analyst ______ Manager ______ K. Smith

This report applies only to the sample investigated and is not necessarily indicative of the quality of apparently identical or similar samples. The liability of the laboratory is limited to the amount paid for the report by the issuee. The issuee assumes all liability for the further distribution of this report or its contents and by making such distribution agrees to hold the laboratory harmless against all claims of persons so informed of the contents hereof.

Lab-3 1/86

Laucks Testing Laboratories, Inc.

Certificate

LABORATORY NO. 7763

DATE: Feb. 2, 1988

940 South Harney St.. Seattle. Washington 98108 (206)767-5060

Chemistry Microbiology, and Technical Services

CLIENT: Kennedy Jenks Chilton Consulting Engineers

33301 Ninth Ave. S., Suite 100

Federal Way, WA 98003

ATTN: Dana Grant

REPORT ON: MATERIAL

(Tan solids)

SAMPLE

IDENTIFICATION: Submitted 1/13/88

Sample was analyzed qualitatively and semi-quantitatively for the purpose of a waste profile sheet.

Note: Large chunks of the sample were ground with a mortar and pestal in order to achieve a more uniform composite sample for analysis.

Physical Characteristics

Color	solid single-phased
1:1 DIW slurry	5.0
Odor	phenolic
Debris	
Total Solids, % by weight	90.5
Viscosity	N/A
Specific gravity @ 25 degrees C	N/A
Flash point	>200°F







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Chemistry Microbiology, and Technical Services

PAGE NO. 2

LABORATORY NO. 7763

Kennedy Jenks Chilton

Chemical Composition

The material was a single-phased large chunky black solid with 11 ppm phenol.

Total Solids @ 105°C ----- 90.5%

Moisture and Volatile Matter @ 105°C ----- 9.5%

E.P. Tox Metals

parts per million

Arsenic Barium Cadmium Chromium Mercury Lead Selenium Silver Copper Nickel Zinc	<0.2 <0.1 <0.01 <0.1 <0.005 <0.1 <0.2 <0.1 <0.1 <0.1
Thallium	<2.







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PAGE NO. 3

LABORATORY NO. 7763

Kennedy Jenks Chilton

Other Components

Total basis, parts per million

Cyanides	<10.
Phenolics	11.
Amines	N/A
PCBs	N/A
Sulfides	<20.

Hazardous Characteristics

د ګ

None Reactive

Dangerous Waste Criteria

Sample was not examined.







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Chemistry Microbiology, and Technical Services

PAGE NO. 4

LABORATORY NO. 7763

Kennedy Jenks Chilton

<u>Comments</u>

Some of the above observations were based on information submitted by our client and by physical observations only.

If you have reason to believe other components or hazardous characteristics are present, appropriate changes should be made when filling out the waste manifest sheet.

When filling out the manifest sheet, any value here shown as N/A (not applicable), should be recorded as a zero or as less than some value if you are confident there is no source for the item.

Key

< = less than

> = greater than

Respectfully submitted,

Laucks Testing Laboratories, Inc.

Therese Lombard

TL:veg



Certificate

940 South Harney St., Seattle, Washington 98108 (206)767-5060

Chemistry Microbiology and Technical Services

CLIENT: Kennedy Jenks Chilton Consulting Engineers

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003

ATTN: Dana Grant

LABORATORY NO. 8185

DATE: Feb. 29, 1988

Job No. 866738

REPORT ON: MATERIAL

SAMPLE

IDENTIFICATION: Submitted 02/05/88 and identified as shown below:

TIP B-5-9 11/19/87 (Drum soluts)

TESTS PERFORMED AND RESULTS:

The sample was analyzed using GC-Ion Trap methods of analysis for solvent identification with results as follows:

F001 - F005 Solvents	%
Acetone n-Butyl alcohol Carbon Disulfide Carbontetrachloride Chlorobenzene Cresols (and cresylic acid) Cyclohexanone 1,2-Dichlorobenzene Ethyl acetate Ethyl benzene Ethyl ether Isobutanol	<0.02 <0.05 <0.02 <0.02 <0.02 <0.1 <0.2 <0.02 <0.02 <0.02 <0.02 <0.02





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PAGE NO. 2

LABORATORY NO. 8185

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F001 - F005 Solvents	%
Methanol	<0.2
Methylene chloride	<0.02
Methyl ethyl ketone	<0.02
Methyl isobutyl ketone	<0.04
Nitrobenzene	<0.04
Pyridine	<0.04
Tetrachloroethylene	<0.02
Toluene	<0.01
1,1,1-Trichloroethane	<0.03
1,1,2-Trichlorotrifluoroethane	<0.02
Trichloroethylene	<0.01
Trichlorofluoromethane	<0.01
Xylene	<0.01

Total F List Solvents ----- Less than 1.0%

<u>Solvents</u>	%
Benzene	<0.1
2-Butanone	<0.1
n-Butyl acetate	<0.1
Butyl cellosolve	<0.1
Cellosolve acetate	<0.1
Cellosolve(ethyl)	<0.5
Chloroform	<0.1
Cyclohexane	<0.1





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Chemistry Microbiology, and Technical Services

PAGE NO. 3

LABORATORY NO. 8185

Kennedy Jenks Chilton Consulting Engineers

<u>Solvents</u>	<u>%</u>
Di-iconyonyl othou	.0.0
Di-isopropyl ether	<0.2
Diacetone alcohol	<0.1
1,4-Dichlorobenzene	<0.1
1,2-Dichloroethane	<0.1
t-1,2-Dichloroethylene	<0.2
1,2-Diethylbenzene	<0.1
1,4-Dioxane	<0.1
2(2ethoxyethoxy)ethanol)	<2.
Ethyl formate	<0.1
n-Hexane	<0.1
2-Hexanone	<0.2
Iso-octane	<0.1
Isobutyl acetate	<0.1
Isopropanol	<0.1
Isopropyl acetate	<0.1
Mesitylene oxide	<0.1
(2(2Methoxyethoxy)ethanol)	<2.
Methyl cellosolve	<1.
2-Nitropropane	<0.1
2,4-Pentadione	<0.1
Phenyl cellosolve	<1.
n-Propanol	<0.5
Styrene	<0.1
1,1,2,2-Tetrachloroethane	<0.1
Tetrahydrofuran	0.03
Tetramethylbenzenes	<0.1
1,1,2-Trichloroethane	<0.1
Trimethyl benzenes	<0.1





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Chemistry Microbiology, and Technical Services

PAGE NO. 4

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 8185

Sample analyzed for Gravimetric Polycyclic Aromatic Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303. The method requires analysis of the samples through successive stages until the result obtained is less than 1% by weight (as received basis) or until the fourth stage has been completed. Results are as shown below:

St	aa	e
Ju	ч	•

% by weight, as received basis

1: Soxhlet Extraction	11.2	•	
Acid-base 2: clean up	8.08		
Silica gel 3: chromatography	6.91	1:	est of the Control of
High Performance 4: Liquid Chromatography (HPLC) Analysis	3.65		

Comment

The reported results for the 4-, 5-, and 6-ring PNAs are based on a correction of the results of the silica gel fractionation for the concentrations of the 2- and 3-ring PNAs in the residue from the silica gel fractionation. The concentrations of the 2- and 3-ring PNAs were determined by HPLC. This calculation is performed in accordance with the method as written.

A qualitative review of the elution region for the 4-, 5-, and 6-ring PNAs in the HPLC chromatogram indicates that the correction factor derived as previously described does not accurately quantify the 4-, 5-, and 6-ring PNAs in this sample. Further confirmatory work, such as GC/MS, may be required to more accurately quantify these compounds in this sample.





Certificate

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Chemistry, Microbiology, and Technical Services

Kennedy Jenks Chilton Consulting Engineers

PAGE NO. 5

LABORATORY NO. 8185

Kev

< indicates "less than"

Respectfully submitted,

Laucks Testing Laboratories, Inc.

J. M. Owens

JMO:emt



Laucks Testing Laboratories, Inc. 940 South Harney St. Seattle. Washington 98108 (206):767-5060



Certificate

Chemistry Microbiology, and Technical Services

APPENDIX

Copy of Chain-of-Custody is Attached



Laucks Testing Laboratories, Inc. 940 South Harney St. Seattle. Washington 98108 (206)767-5060



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Chemistry Microbiology and Technical Services

PAGE NO. 11

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 5

5

Ignitablity

Note A

Note A = Does not flash at, or less than 200°F, when tested in a Setaflash tester (and does not burn when exposed to open flame).

Sample No. 5 was analyzed for E.P. Toxicity in accordance with <u>Test Methods for Evaluating Solid Waste</u>, (SW 846), U.S.E.P.A., November, 1986. Extraction was performed using Method 1310. Mercury was determined using Method 7470; other metals performed by ICAP, Method 6010.

parts per million (mg/L)

	5	MCL
Arsenic	L/0.2	5.0
Barium	0.6	100.
Cadmium	L/0.01	1.0
Chromium	L/0.1	5.0
Lead	L/0.1	5.0
Mercury	L/0.005	0.2
Selenium	L/0.2	1.0
Silver	L/0.1	5.0

Comment:

Due to insufficient sample, less than the required 100 gm weight was extracted, proportionally.

Sample size extracted: 17.3g



This report is submitted for the exclusive use of the person, partnership, or corporation to whom it is addressed. Subsequent use of the name of this company or any member of its staff in connection with the advertising or sale of any product or process will be granted only on contract. This company accepts no responsibility except for the due performance of inspection and/or analysis in good faith and according to the rules of the trade and of science.

Laucks Testing Laboratories, Inc. 940 South Harney St., Seattle. Washington 98108 (206)767-5060



Certificate

Chemistry Microbiology and Technical Services

PAGE NO. 12

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 6

6

Total Solids, %

94.5

parts per million (mg/kg), dry basis

011 & Grease

5.700.

Sample No. 6 was analyzed for Halogenated Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303 with results as follows:

parts per million (mg/kg), as received basis

_6

Halogenated Hydrocarbons*

44.

*Reported as the sum of the halogens bromide, chloride fluoride and iodide. A value of less than 100 mg/kg is classified as undesignated waste.





940 South Harney St., Seattle, Washington 98108 (206)767-5060

Chemistry, Microbiology, and Technical Services

PAGE NO. 13

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 6

6

Ignitablity

Note A

Note A = Does not flash at, or less than 200°F, when tested in a Setaflash tester (and does not burn when exposed to open flame).

Sample No. 6 was analyzed for E.P. Toxicity in accordance with <u>Test Methods for Evaluating Solid Waste</u>, (SW 846), U.S.E.P.A., November, 1986. Extraction was performed using Method 1310. Mercury was determined using Method 7470; other metals performed by ICAP, Method 6010.

parts per million (mg/L)

	<u>_6</u> _	<u>MCL</u>
Arsenic Barium Cadmium	L/0.2 1.4 0.06	5.0 100. 1.0
Chromium	L/0.1	5.0
Lead Mercury	0.3 L/0.005	5.0 0.2
Selenium	L/0.2	1.0
Silver	L/0.1	5.0

Comment:

Due to insufficient sample, less than the required 100 gm weight was extracted, proportionally.

Sample size extracted: 17.8 g





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Chemistry Microbiology and Technical Services

PAGE NO. 14

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Comments:

Some of the above observations were based on information submitted by our client and physical observations only.

If you have reason to believe other components or hazardous characteristics are present, appropriate changes should be made when filling out the waste manifest sheet.

When filling out the manifest sheet, any value here shown as N/A (not applicable), should be recorded as a zero or as less than some value if you are confident there is no source for the item.

<u>Key</u>

< indicates "less than"
MCL = Maximum Contamination Level allowed per regulation.</pre>

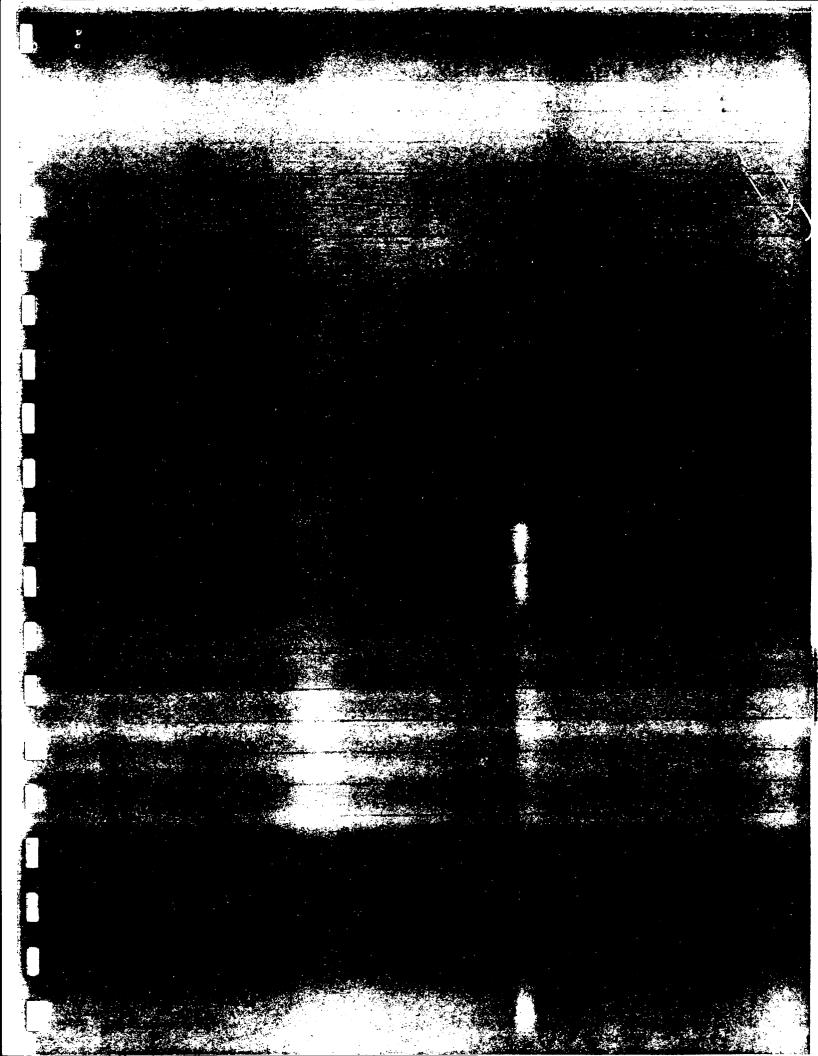
Respectfully submitted.

Laucks Testing Laboratories, Inc.

J. M. Owens

JMO: laj





CHEMICAL PROCESSORS, INC./RESOURCE RECOVERY CORP.

2203 Airport Way So., Suite 400 ● Seattle, WA 98134 Chempro 223-0500 ● Resource Recovery 223-0900

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		P. Designated Facility Name and Site Address 10. Chempra 734 So. Lucile St. Seattle, WA (206) 767-3362 □ Chempra Fier 91 Seattle, WA (206) 284-2450 □ Chempra 1701 Alexander Tacoma, WA (206) 627-7568	WAD 000812909 WAD 000812917 WAD 020257945			the de the second	<u> </u>	
	L	☐ McClary/Columbia 625 So. 32nd St. Washougal, WA (206)835-8594 ☐ Other:	WAD 092300250		<u> </u>	76	, 	3362
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	١	If I am a large quantity generator, I certify that I have a program in place to re determined to be economically practicable and that I have selected the practical						
		which minimizes the present and future threat to human health and the environm effort to minimize my waste generation and select the best waste management met	ent, OR, if I om a small qua	ntity genero	tor, I hov	e made a good fait	h	
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CHEMICAL PROCESSORS, INC./RESOURCE RECOVERY CORP. 2203 Airport Wey So., Suite 400 • Seattle, WA 98134 Chempro 223-0500 • Resource Recovery 223-0900

	lease print or type form designed for use on elite (12-pitch) typewriter.)				Form Approved OM/	B No. 2050)-0039 Expires 9:30.8
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CHEMICAL PROCESSORS, INC./RESOURCE RECOVERY CORP.

2203 Airport Way So., Suite 400 ● Seattle, WA 98134 Chempro 223-0500 ● Resource Recovery 223-0900

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CHEMICAL PROCESSORS, INC./RESOURCE RECOVERY CORP.

2203 Airport Way So., Suite 400 ● Seattle, WA 98134 Chempro 223-0500 ● Resource Recovery 223-0900

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Kennedy/Jenks/Chilton Laboratory Divis

Field Sample Chain of Custody Record

Source of Sample(s) TIP	·	Collect	or Dana Grant
Address		Affilia	cion KJ/C
		Address	33301 9+HAJS SUITE
Phone ()	·		Federal Wy, WA 9800
Report to (1) Dana Grant		Phone	(206) 874-0555
Sample Information			K/J/C Job Number <u>866738</u>
Lab No. Field No. Date Time	Type (2)	Flow Rate	Remarks (Suspected Contaminants, Field Conditions, etc.)
B-5-9 11	solid		PAH
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2.			
Logged in at K/J/C by			
2065EG TH BC 2/0/0 07			

(2) e.g. water, sludge, soil, etc.

⁽¹⁾ There is a separate Request for Analysis form that should be filled out by the colland given to the Laboratory when samples are delivered.

⁽³⁾ If any samples are not intact at time of transfer, please describe on the back of t form.

Testing Laboratories, Inc.



Certificate

LABORATORY NO. 7225

DATE: Jan. 4, 1988

Committee of the state of the s

940 South Harney St., Seattle, Washington 98108 (206)767-5060

Chemistry Microbiology and Technical Services

CLIENT: Kennedy Jenks Chilton Consulting Engineers

33301 Ninth Avenue South, Suite 100

Federal Way, WA 98003 ATTN: Nathan Graves

Job No. 866738

REPORT ON:

IDENTIFICATION:

SAMPLE

LIQUIDS AND SOLIDS

Submitted 12/03/87 and identified as shown below:

1) Composite 12/01/87 (liquid) 2) B-4 12/02/87 (acid) 3) BT-3 12/02/87 (sludge)

4) Composite Solids 12/02/87 (solids) 12/02/87

(solids) 5) C-10 . 6) C-9 12/02/87 (solids)

TESTS PERFORMED AND RESULTS:

Sample No. 1

Sample No. 1 was analyzed for Halogenated Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303 with results as follows:

parts per million (mg/kg), as received basis

Halogenated Hydrocarbons*

.

38.

*Reported as the sum of the halogens bromide, chloride fluoride and iodide. A value of less than 100 mg/kg is classified as undesignated waste.





Certificate

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Chemistry Microbiology and Technical Services

PAGE NO. 2

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 1

Sample No. 1 was analyzed qualitatively and semi-quantitatively for the purpose of classification for a waste profile sheet with observations and results as follow:

Physical Characteristics

Color	Bi-Layered Top: Black
Physical State at 70 degrees FLayers	Bottom: Light Brown Liquid Bi-Layered
Free Liquid	Top: 15% Bottom: 85% 98.+% 7.5
Odor Debris Total Solids (by weight) Viscosity	Oil/Solvent Sediment 5.7% Top: Oil
Specific GravityFlash Point	Bottom: Water 0.91 101-139°F

Chemical Composition

The material was bi-layered with a top black oil and solvent phase (15% of the total by volume) and a bottom light brown aqueous phase (85% of the total by volume).

Gas chromatographic analysis by ion trap and total solids were performed with results as follows:

Total Solids at 105°C	5.7%	
Toluene	0.1%	
Xylene	0.6%	
1,3,5-Trimethy Thenzene	1.5%	
1,3-Diethylbenzene	0.1%	
1,2,4-Trimethylbenzene	4.8%	
Other Alkyl Benzenes	6.0%	(approximately)
Moisture (by difference)	81.2%	



This report is submitted for the exclusive use of the person, partnership, or corporation to whom it is addressed. Subsequent use of the name of this company or any member of its staff in connection with the advertising or sale of any product or process will be granted only on contract. This company accepts no responsibility except for the due performance of inspection and/or analysis in good faith and according to the rules of the trade and of science.





Certificate

Chemistry, Microbiology, and Technical Services

PAGE NO. 3

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 1

Sample 1 was analyzed for E.P. Toxicity in accordance with <u>Test Methods for Evaluating Solid Waste</u>, (SW 846), U.S.E.P.A., November, 1986. Extraction was performed using Method 1310. Mercury was determined using Method 7470; other metals performed by ICAP, Method 6010.

EP Tox Metals	parts per million
Arsenic Barium Cadmium Chromium Mercury Lead Selenium Silver Copper Nickel Zinc Thallium	L/0.2/ 302 and 2.745/3 1.3 L/0.01 L/0.1 L/0.005 0.1 L/0.2 L/0.1 L/0.1 L/0.1
Other Components	
Cyanides Phenols Amines PCBs Sulfides	L/10. L/10. N/A N/A L/20.
Hazardous Characteristics	
None Reactive	
Dangerous Waste Criteria	
Sample Examined	No





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Chemistry, Microbiology, and Technical Services

PAGE NO. 4

LABORATORY NO. 7225

Kennedy Jenks Chilton Consulting Engineers

Sample No. 2

Acidity, calculated as % HCL to methyl orange end point ---- 30.0

Total Metals

parts per million

	<u> 2</u>
Arsenic Bartum	ভাষা - L/5-মালের বিল্লাস ভাষার বিষয়ে ভাল বিশ্ব হল L/2- তালেল বিল্লাস্থান স্থাপন্ত বিল্লাস্থ
Cadmium@diates of the second	is the many and the state of the second
Chromium	L/2.
Mercury	L/5.
Lead	L/2.
Selenium	L./5
Silver	L/2.
Copper	L/2.
Nickel	L/2.
Zinc	L/2.
Thallium	L/50.

Committee of the Commit







940 South Harney St., Seattle Washington 98108 (206)767-5060

Chemistry Microbiology and Technical Services

PAGE NO. 5

Kennedy Jenks Chilton Consulting Engineers

LABORATORY NO. 7225

Sample No. 3

Sample: No.: 3 was analyzed for Halogenated Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303 with results as follows:

parts per million (mg/kg), as received basis

3

Halogenated Hydrocarbons*

à ..

210.

*Reported as the sum of the halogens bromide, chloride fluoride and iodide. A value of 100-10,000 ppm is classified as dangerous waste.

Sample No. 3 was analyzed qualitatively and semi-quantitatively for the purpose of classification for a waste profile sheet with observations and results as follow:

Physical Characteristics

Color	Black .
Physical State at 70 degrees F	Semi-Solid
Layers	Single Phased
Free Liquid	No
pH (1:1 DIW slurry), glass electrode at 25 degrees C	4.2
Odor	Tar/Solvent
Debris	No 47.2%
Total Solids (by weight)	47.2% N/A
Specific Gravity	
Flash Point	.dv/70°F







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Chemistry, Microbiology, and Technical Services

PAGE NO. 6

LABORATORY NO. 7225

Kennedy Jenks Chilton Consulting Engineers

Sample No. 3

Chemical Composition

The material was a single phased black semi-solid with a solvent/tar-like odor.

Gas chormatographic analysis by ion trap and total solids were performed with results as follows:

Total Solids at 105°C	47.2%
Toluene	0.3%
Xylene	0.6%
n-Hexane	1.2%
Cyclohexane	5.3%
Octane	5.6%
Nonane	5.5%
1,3,5-Trimethylbenzene	0.1%
1,2,4-Trimethylbenzene	0.1%
A light hydrocarbon pattern including	
a gasoline pattern (by difference)	34.1%







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Sample No. 3

Sample 3 was analyzed for E.P. Toxicity in accordance with <u>Test Methods for Evaluating Solid Waste</u>, (SW 846), U.S.E.P.A., November, 1986. Extraction was performed using Method 1310. Mercury was determined using Method 7470; other metals performed by ICAP, Method 6010.

EP Tox Metals	parts per million
Arsenic Barium Cadmium Chromium Mercury Lead Selenium	L/0.2 L/0.1 L/0.01 L/0.1 L/0.005 L/0.1 L/0.2
Silver Copper Nickel Zinc Thallium Other Components	L/0.1 L/0.1 L/0.1 L/0.1 L/2.
Cyanides Phenols Amines PCBs Sulfides	L/10. L/10. N/A N/A L/20.
None Reactive	
Dangerous Waste Criteria Sample Examined	No







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Sample No. 4

4

Total Solids. %

97.0

parts per million (mg/kg), dry basis

011 & Grease

340.

Sample No. 4 was analyzed for Halogenated Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303 with results as follows:

parts per million (mg/kg), as received basis

4

Halogenated Hydrocarbons*

15.

*Reported as the sum of the halogens bromide, chloride fluoride and iodide. A value of less than 100 mg/kg is classified as undesignated waste.





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Sample No. 4

4

Ignitablity

Note A

Note A = Does not flash at, or less than 200°F, when tested in a Setaflash tester (and does not burn when exposed to open flame).

Sample No. 4 was analyzed for E.P. Toxicity in accordance with <u>Test Methods for Evaluating Solid Waste</u>, (SW 846), U.S.E.P.A., November, 1986. Extraction was performed using Method 1310. Mercury was determined using Method 7470; other metals performed by ICAP, Method 6010.

parts per million (mg/L)

·		MCL
Arsenic	L/0.2	5.0
Barium	0.2	100.
Cadmium	L/0.01	1.0
Chromium	L/0.1	5.0
Lead	L/0.1	5.0
Mercury	L/0.005	0.2
Selenium	L/0.2	1.0
Silver	L/0.1	5.0

Comment:

Due to insufficient sample, less than the required 100 gm weight was extracted, proportionally.

Sample size extracted: 94.3 g



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Sample No. 5

5

Total Solids. %

77.6

parts per million (mg/kg), dry basis

011 & Grease

180.

Sample No. 5 was analyzed for Halogenated Hydrocarbons in accordance with Washington State Department of Ecology WAC 173-303 with results as follows:

parts per million (mg/kg), as received basis

5

Halogenated Hydrocarbons*

30.

*Reported as the sum of the halogens bromide, chloride fluoride and iodide.

A value of less than 100 mg/kg is classified as undesignated waste.

